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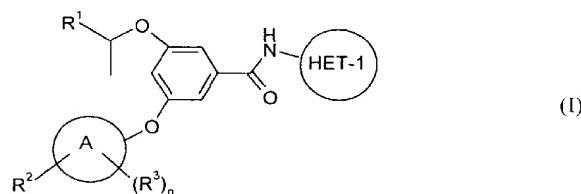
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(54) Title: BENZOYL AMINO HETEROCYCLYL COMPOUNDS AS GLUCOKINASE (GLK) ACTIVATORS



(57) Abstract: The invention relates to a novel group of compounds of Formula (I) or a salt thereof: wherein R¹, R², R³, n, A and HET-1 are as described in the specification, which may be useful in the treatment or prevention of a disease or medical condition mediated through glucokinase (GLK) such as type 2 diabetes. The invention also relates to pharmaceutical compositions comprising said compounds, methods of treatment of diseases mediated by GLK using said compounds and methods for preparing compounds of Formula (I).

WO 2008/050117 A1

BENZOYL AMINO HETEROCYCLYL COMPOUNDS AS GLUCOKINASE (GLK) ACTIVATORS

The present invention relates to a group of benzoyl amino heterocyclyl compounds which are useful in the treatment or prevention of a disease or medical condition mediated through glucokinase (GLK or GK), leading to a decreased glucose threshold for insulin secretion. In addition the compounds are predicted to lower blood glucose by increasing hepatic glucose uptake. Such compounds may have utility in the treatment of Type 2 diabetes and obesity. The invention also relates to pharmaceutical compositions comprising said compounds and to methods of treatment of diseases mediated by GLK using said compounds.

In the pancreatic β -cell and liver parenchymal cells the main plasma membrane glucose transporter is GLUT2. Under physiological glucose concentrations the rate at which GLUT2 transports glucose across the membrane is not rate limiting to the overall rate of glucose uptake in these cells. The rate of glucose uptake is limited by the rate of phosphorylation of glucose to glucose-6-phosphate (G-6-P) which is catalysed by glucokinase (GLK) [1]. GLK has a high (6-10mM) Km for glucose and is not inhibited by physiological concentrations of G-6-P [1]. GLK expression is limited to a few tissues and cell types, most notably pancreatic β -cells and liver cells (hepatocytes) [1]. In these cells GLK activity is rate limiting for glucose utilisation and therefore regulates the extent of glucose induced insulin secretion and hepatic glycogen synthesis. These processes are critical in the maintenance of whole body glucose homeostasis and both are dysfunctional in diabetes [2].

In one sub-type of diabetes, Maturity-Onset Diabetes of the Young Type 2 (MODY-2), the diabetes is caused by GLK loss of function mutations [3, 4].

Hyperglycaemia in MODY-2 patients results from defective glucose utilisation in both the pancreas and liver [5]. Defective glucose utilisation in the pancreas of MODY-2 patients results in a raised threshold for glucose stimulated insulin secretion. Conversely, rare activating mutations of GLK reduce this threshold resulting in familial hyperinsulinism [6, 6a, 7]. In addition to the reduced GLK activity observed in MODY-2 diabetics, hepatic glucokinase activity is also decreased in type 2 diabetics [8]. Importantly, global or liver selective overexpression of GLK prevents or reverses the development of the diabetic phenotype in both dietary and genetic models of the disease [9-12]. Moreover, acute

- 2 -

treatment of type 2 diabetics with fructose improves glucose tolerance through stimulation of hepatic glucose utilisation [13]. This effect is believed to be mediated through a fructose induced increase in cytosolic GLK activity in the hepatocyte by the mechanism described below [13].

5 Hepatic GLK activity is inhibited through association with GLK regulatory protein (GLKRP). The GLK/GLKRP complex is stabilised by fructose-6-phosphate (F6P) binding to the GLKRP and destabilised by displacement of this sugar phosphate by fructose-1-phosphate (F1P). F1P is generated by fructokinase mediated phosphorylation of dietary fructose. Consequently, GLK/GLKRP complex integrity and hepatic GLK activity
10 is regulated in a nutritionally dependent manner as F6P is dominant in the post-absorptive state whereas F1P predominates in the post-prandial state. In contrast to the hepatocyte, the pancreatic β -cell expresses GLK in the absence of GLKRP. Therefore, β -cell GLK activity is regulated extensively by the availability of its substrate, glucose. Small molecules may activate GLK either directly or through destabilising the GLK/GLKRP
15 complex. The former class of compounds are predicted to stimulate glucose utilisation in both the liver and the pancreas whereas the latter are predicted to act selectively in the liver. However, compounds with either profile are predicted to be of therapeutic benefit in treating Type 2 diabetes as this disease is characterised by defective glucose utilisation in both tissues.

20 GLK, GLKRP and the K_{ATP} channel are expressed in neurones of the hypothalamus, a region of the brain that is important in the regulation of energy balance and the control of food intake [14-18]. These neurones have been shown to express orectic and anorectic neuropeptides [15, 19, 20] and have been assumed to be the glucose-sensing neurones within the hypothalamus that are either inhibited or excited by changes in ambient glucose concentrations [17, 19, 21, 22]. The ability of these neurones to sense changes in glucose levels is defective in a variety of genetic and experimentally induced models of obesity [23-28]. Intracerebroventricular (icv) infusion of glucose analogues, that are competitive inhibitors of glucokinase, stimulate food intake in lean rats [29, 30]. In contrast, icv infusion of glucose suppresses feeding [31]. Thus, small molecule
25 activators of GLK may decrease food intake and weight gain through central effects on GLK. Therefore, GLK activators may be of therapeutic use in treating eating disorders, including obesity, in addition to diabetes. The hypothalamic effects will be additive or
30

- 3 -

synergistic to the effects of the same compounds acting in the liver and/or pancreas in normalising glucose homeostasis, for the treatment of Type 2 diabetes. Thus the GLK/GLKRP system can be described as a potential "Diabesity" target (of benefit in both Diabetes and Obesity).

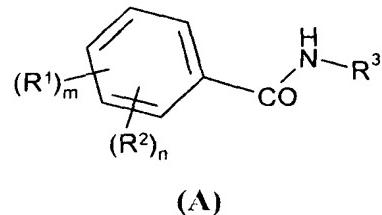
GLK is also expressed in specific entero-endocrine cells where it is believed to control the glucose sensitive secretion of the incretin peptides GIP (glucose-dependent insulinotropic polypeptide) and GLP-1 (Glucagon-Like Peptide-1) from gut K-cells and L-cells respectively (32, 33, 34). Therefore, small molecule activators of GLK may have additional beneficial effects on insulin secretion, b-cell function and survival and body weight as a consequence of stimulating GIP and GLP-1 secretion from these entero-endocrine cells.

In WO00/58293 and WO01/44216 (Roche), a series of benzylcarbamoyl compounds are described as glucokinase activators. The mechanism by which such compounds activate GLK is assessed by measuring the direct effect of such compounds in an assay in which GLK activity is linked to NADH production, which in turn is measured optically - see details of the *in vitro* assay described hereinafter. Compounds of the present invention may activate GLK directly or may activate GLK by inhibiting the interaction of GLKRP with GLK.

Further GLK activators have been described in WO03/095438 (substituted phenylacetamides, Roche), WO03/055482 (carboxamide and sulphonamide derivatives, Novo Nordisk), WO2004/002481 (arylcarbonyl derivatives, Novo Nordisk), and in WO03/080585 (amino-substituted benzoylaminoheterocycles, Banyu).

Our International application Number: WO03/000267 describes a group of benzoyl amino pyridyl carboxylic acids which are activators of the enzyme glucokinase (GLK).

Our International application Number: WO03/015774 describes compounds of the Formula (A):



- 4 -

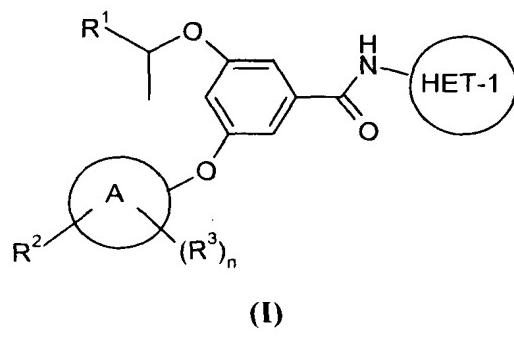
wherein R³ is phenyl or a substituted heterocycle other than a carboxylic acid substituted pyridyl.

International application WO2004/076420 (Banyu) describes compounds which are generally a subset of those described in WO03/015774, wherein for example R¹ is an (substituted) alkyl ether and R² is (substituted) phenoxy.

We have surprisingly found a small group of compounds, generally a selected subgroup of those described in WO 03/015774, which have generally superior potency for the GLK enzyme, and more advantageous physical properties, including, for example, higher aqueous solubility, higher permeability, and/or lower plasma protein binding.

Consequently, such compounds having a balance of these properties would be expected to display higher plasma free drug levels and superior in vivo efficacy after oral dosing as determined, for example, by activity in Oral Glucose Tolerance Tests (OGTTs). Therefore this group of compounds would be expected to provide superior oral exposure at a lower dose and thereby be particularly suitable for use in the treatment or prevention of a disease or medical condition mediated through GLK. The compounds of the invention may also have superior potency and/or advantageous physical properties (as described above) and/or favourable toxicity profiles and/or favourable metabolic profiles in comparison with other GLK activators known in the art, as well as those described in WO 03/015774.

Thus, according to the first aspect of the invention there is provided a compound of Formula (I):



wherein:

R¹ is selected from fluoromethoxymethyl, difluoromethoxymethyl and

trifluoromethoxymethyl;

R² is a substituent on a carbon atom of Ring A and is selected from -C(O)NR⁴R⁵,

-SO₂NR⁴R⁵, -S(O)_pR⁴ and HET-2;

- 5 -

HET-1 is a 5- or 6-membered, C-linked heteroaryl ring containing a nitrogen atom in the 2-position relative to the amide nitrogen to which the ring is attached and optionally 1 or 2 further ring heteroatoms independently selected from O, N and S; which ring is optionally substituted on an available carbon atom, or on a ring nitrogen atom provided it is not thereby quaternised, with 1 or 2 substituents independently selected from R⁶;

HET-2 is a 4-, 5- or 6-membered, C- or N-linked heterocycll ring containing 1, 2, 3 or 4 heteroatoms independently selected from O, N and S, wherein a -CH₂- group can optionally be replaced by a -C(O)-, and wherein a sulphur atom in the heterocyclic ring may optionally be oxidised to a S(O) or S(O)₂ group, which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁷;

R³ is a substituent on a carbon atom of Ring A and is selected from halo;

R⁴ is selected from hydrogen, (1-4C)alkyl [optionally substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and -C(O)NR⁵R⁵], (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and HET-2;

R⁵ is hydrogen or (1-4C)alkyl;

R⁶ is independently selected from (1-4C)alkyl, hydroxy(1-4C)alkyl, (1-4C)alkoxy(1-4C)alkyl, (1-4C)alkylS(O)p(1-4C)alkyl, amino(1-4C)alkyl, (1-4C)alkylamino(1-4C)alkyl, di(1-4C)alkylamino(1-4C)alkyl, and/or (for R⁶ as a substituent on carbon) halo;

R⁷ is selected from (1-4C)alkyl, -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, (1-4C)alkoxy(1-4C)alkyl, hydroxy(1-4C)alkyl, -S(O)pR⁵ and/or (for R⁷ as a substituent on carbon) hydroxy and (1-4C)alkoxy;

Ring A is a 5- or 6- membered heteroaryl ring, containing 1, 2 or 3 ring heteroatoms independently selected from O, S and N; which ring optionally further substituted on an available nitrogen atom (provided it is not thereby quaternised) by a substituent selected from R⁸;

R⁸ is selected from (1-4C)alkyl, (3-6C)cycloalkyl, hydroxy(1-4C)alkyl, (1-4C)alkoxy(1-4C)alkyl, -C(O)(1-4C)alkyl, benzyl, and (1-4C)alkylsulfonyl;

p is (independently at each occurrence) 0, 1 or 2;

n is 0, 1 or 2;

or a salt thereof.

It will be understood that when R⁴ is (1-4C)alkyl substituted with -C(O)NR⁵R⁵, each R⁵ is independently selected from hydrogen and (1-4C)alkyl, and therefore this definition of R⁴ includes (but is not limited to) (1-4C)alkyl substituted with -CONH₂, -CONHMe, -CONMe₂ or -CONMeEt.

5 It will be understood that where a compound of the formula (I) contains more than one HET-2 ring, they may be the same or different.

It will be understood that where a compound of the formula (I) contains more than one group R⁴, they may be the same or different.

10 It will be understood that where a compound of the formula (I) contains more than one group R⁵, they may be the same or different.

It will be understood that where a compound of the formula (I) contains more than one group R³, they may be the same or different.

A similar convention applies for all other groups and substituents on a compound of formula (I) as hereinbefore defined.

15 Compounds of Formula (I) may form salts which are within the ambit of the invention. Pharmaceutically-acceptable salts are preferred although other salts may be useful in, for example, isolating or purifying compounds.

In another aspect, the invention relates to compounds of formula (I) as hereinabove defined or to a pharmaceutically-acceptable salt.

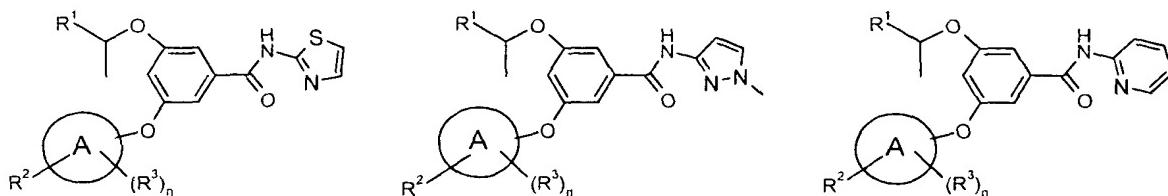
20 In another aspect, the invention relates to compounds of formula (I) as hereinabove defined or to a pro-drug thereof. Suitable examples of pro-drugs of compounds of formula (I) are in-vivo hydrolysable esters of compounds of formula (I). Therefore in another aspect, the invention relates to compounds of formula (I) as hereinabove defined or to an in-vivo hydrolysable ester thereof.

25 In this specification the generic term "alkyl" includes both straight-chain and branched-chain alkyl groups. However references to individual alkyl groups such as "propyl" are specific for the straight chain version only and references to individual branched-chain alkyl groups such as *t*-butyl are specific for the branched chain version only. For example, "(1-4C)alkyl" includes methyl, ethyl, propyl, isopropyl and *t*-butyl. An analogous convention applies to other generic terms.

30 For the avoidance of doubt, reference to the group HET-1 containing a nitrogen in the 2-position, is intended to refer to the 2-position relative to the amide nitrogen atom to

- 7 -

which the group is attached. For example, the definition of formula (I) encompasses (but is not limited to) the following structures:



5 Suitable examples of HET-1 as a 5- or 6-membered, C-linked heteroaryl ring as hereinbefore defined, include thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl, pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl and triazolyl.

10 It will be understood that HET-2 can be a saturated, or partially or fully unsaturated ring.

Suitable examples of HET-2 include azetidinyl, furyl, thienyl, thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl, pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl, morpholino, morpholinyl, piperidinyl, piperazinyl, morpholinyl, thiomorpholinyl, pyrrolyl, pyrrolidinyl, pyrrolidonyl, 15 2,5-dioxopyrrolidinyl, 1,1-dioxotetrahydrothienyl, 2-oximidazolidinyl, 2,4-dioxoimidazolidinyl, 2-oxo-1,3,4-(4-triazolinyl), 2-oxazolidinonyl, 2-oxotetrahydrofuranyl, tetrahydrofuranyl, tetrahydropyranyl, 1,1-dioxothiomorpholino, 1,3-dioxolanyl, 1,2,4-triazolyl, 1,2,3-triazolyl, pyranyl, and 4-pyridonyl.

20 It will be understood that HET-2 may be linked by any appropriate available C or N atom, therefore for example, for HET-2 as "imidazolyl" includes 1-, 2-, 4- and 5-imidazolyl.

Suitable examples of Ring A as hereinbefore defined include thienyl, furyl, thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl, pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl and triazolyl. Further suitable examples of 25 Ring A include aromatic heterocycles where a ring nitrogen or sulfur atom has been oxidised but aromaticity has been preserved, for example a pyridine N-oxide. Further suitable examples of Ring A include thiazolyl, pyridyl, pyrazinyl, pyridazinyl and pyrimidinyl.

It will be appreciated that the above suitable values for HET-1, HET-2 and Ring A may all be optionally substituted as defined hereinbefore.

It will be appreciated that, where definitions of heterocyclyl groups HET-1, HET-2 and Ring A encompass heteroaryl or heterocyclyl rings which may be substituted on nitrogen, such substitution may not result in charged quaternary nitrogen atoms or unstable structures (such as N-halo compounds). It will be appreciated that the definitions of HET-1, HET-2 and Ring A are not intended to include any O-O, O-S or S-S bonds. It will be appreciated that the definitions of HET-1, HET-2 and Ring A are not intended to include unstable structures.

Examples of **(1-4C)alkyl** include methyl, ethyl, propyl, isopropyl, butyl and tert-butyl; examples of **(1-4C)alkoxy** include methoxy, ethoxy, propoxy, isopropoxy and tertbutoxy; examples of **(3-6C)cycloalkyl** include cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl; examples of **halo** include fluoro, chloro, bromo and iodo; examples of **hydroxy(1-4C)alkyl** include hydroxymethyl, 1-hydroxyethyl, 2-hydroxyethyl, 2-hydroxypropyl, 3-hydroxypropyl, 1-hydroxyisopropyl and 4-hydroxybutyl; examples of **(1-4C)alkoxy(1-4C)alkyl** include methoxymethyl, ethoxymethyl, tert-butoxymethyl, 2-methoxyethyl, 2-ethoxyethyl, methoxypropyl, 2-methoxypropyl and methoxybutyl; examples of **(1-4C)alkylS(O)p(1-4C)alkyl** include methylsulfinylmethyl, ethylsulfinylmethyl, ethylsulfinylethyl, methylsulfinylpropyl, methylsulfinylbutyl, methylsulfonylmethyl, ethylsulfonylmethyl, ethylsulfonylethyl, methylsulfonylpropyl, methylsulfonylbutyl, methylthiomethyl, ethylthiomethyl, ethylthioethyl, methylthiopropyl, and methylthiobutyl; examples of **amino(1-4C)alkyl** include aminomethyl, aminoethyl, 2-aminopropyl, 3-aminopropyl, 1-aminoisopropyl and 4-aminobutyl; examples of **(1-4C)alkylamino(1-4C)alkyl** include (N-methyl)aminomethyl, (N-ethyl)aminomethyl, 1-((N-methyl)amino)ethyl, 2-((N-methyl)amino)ethyl, (N-ethyl)aminoethyl, (N-methyl)aminopropyl, and 4-((N-methyl)amino)butyl; examples of **di(1-4C)alkylamino(1-4C)alkyl** include dimethylaminomethyl, methyl(ethyl)aminomethyl, methyl(ethyl)aminoethyl, (N,N-diethyl)aminoethyl, (N,N-dimethyl)aminopropyl and (N,N-dimethyl)aminobutyl; examples of **(1-4C)alkylamino** include methylamino, ethylamino, propylamino, isopropylamino, butylamino and tert-butyldiamino; examples of **di(1-4C)alkylamino** include dimethylamino, methyl(ethyl)amino, diethylamino, dipropylamino, di-isopropylamino and dibutylamino; examples of **-C(O)(1-4C)alkyl**

include methylcarbonyl, ethylcarbonyl, propylcarbonyl and tert-butyl carbonyl; examples of **(1-4C)alkylsulfonyl** include methylsulfonyl, ethylsulfonyl, isopropylsulfonyl and tert-butylsulfonyl.

It is to be understood that, insofar as certain of the compounds of Formula (I) defined above may exist in optically active or racemic forms by virtue of one or more asymmetric carbon atoms, the invention includes in its definition any such optically active or racemic form which possesses the property of stimulating GLK directly or inhibiting the GLK/GLKRP interaction. The synthesis of optically active forms may be carried out by standard techniques of organic chemistry well known in the art, for example by synthesis from optically active starting materials or by resolution of a racemic form. It is also to be understood that certain compounds may exist in tautomeric forms and that the invention also relates to any and all tautomeric forms of the compounds of the invention which activate GLK.

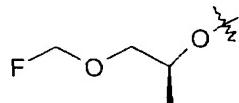
It is also to be understood that certain compounds of the formula (I) and salts thereof can exist in solvated as well as unsolvated forms such as, for example, hydrated forms. It is to be understood that the invention encompasses all such solvated forms which activate GLK.

In one embodiment of the invention are provided compounds of formula (I), in an alternative embodiment are provided pharmaceutically-acceptable salts of compounds of formula (I), in a further alternative embodiment are provided in-vivo hydrolysable esters of compounds of formula (I), and in a further alternative embodiment are provided pharmaceutically-acceptable salts of in-vivo hydrolysable esters of compounds of formula (I).

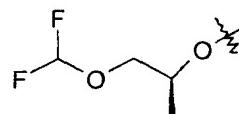
Preferred values of each variable group are as follows. Such values may be used where appropriate with any of the values, definitions, claims, aspects or embodiments defined hereinbefore or hereinafter. In particular, each may be used as an individual limitation on the broadest definition of formula (I). Further, each of the following values may be used in combination with one or more of the other following values to limit the broadest defintion of formula (I).

- 30 (1) R¹ is fluoromethoxymethyl or difluoromethoxymethyl
- (2) R¹ is fluoromethoxymethyl and the configuration is preferably (S), that is the sidechain is:

- 10 -



(3) R¹ is difluoromethoxymethyl and the configuration is preferably (S), that is the sidechain is:



- 5 (4) R² is -C(O)NR⁴R⁵
- (5) R² is -SO₂NR⁴R⁵
- (6) R² is -S(O)_pR⁴
- (7) R² is HET-2
- (8) R² is -C(O)NR⁴R⁵ or -SO₂NR⁴R⁵
- 10 (9) R² is in the para position relative to the ether linkage
- (10) n is 0 or 1
- (11) n is 0
- (12) n is 1, R² is in the para position relative to the ether linkage, R³ is in the ortho position relative to the ether linkage
- 15 (13) n is 1, R² is in the para position relative to the ether linkage, R³ is in the meta position relative to the ether linkage
- (14) n is 1
- (15) n is 2
- (16) n is 2 and both R³ are halo
- 20 (17) n is 2 and each R³ is independently fluoro or chloro
- (18) n is 2, R² is in the para position relative to the ether linkage and each R³ is in an ortho position relative to the ether linkage
- (19) n is 2, both R³ are halo, R² is in the para position relative to the ether linkage and each R³ is in an ortho position relative to the ether linkage
- 25 (20) n is 2, both R³ are halo, R² is in the para position relative to the ether linkage and one R³ is in an ortho position relative to the ether linkage and the other R³ is in a meta position relative to the ether linkage
- (21) R³ is chloro or fluoro

- 11 -

(22) R³ is fluoro

(23) R³ is chloro

(24) n is 2 and both R³ are fluoro

(25) n is 2 and one R³ is fluoro and the other is chloro

5 (26) p is 0

(27) p is 1

(28) p is 2

(29) HET-1 is a 5-membered heteroaryl ring

(30) HET-1 is a 6-membered heteroaryl ring

10 (31) HET-1 is substituted with 1 or 2 substituents independently selected from R⁶

(32) HET-1 is substituted with 1 substituent selected from R⁶

(33) HET-1 is unsubstituted

(34) HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl, pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl, and triazolyl

15 (35) HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, imidazolyl, oxazolyl, isoxazolyl and oxadiazolyl

(36) HET-1 is selected from pyridyl, pyrazinyl, pyridazinyl and pyrimidinyl

(37) HET-1 is optionally substituted pyrazolyl, for example pyrazolyl or N-methylpyrazolyl

20 (38) HET-1 is pyridyl or pyrazinyl

(39) HET-1 is pyrazinyl

(40) HET-1 is selected from pyrazolyl, N-methylpyrazolyl and methylpyrazinyl (such as 5-methylpyrazin-2-yl)

25 (41) HET-1 is pyrazolyl (optionally substituted with ethyl, isopropyl or 1 or 2 methyl), thiazolyl (optionally substituted with methyl), pyrazinyl (optionally substituted with methyl), pyridyl (optionally substituted by fluoro), isoxazolyl (optionally substituted with methyl) and thiadiazolyl (optionally substituted with methyl)

(42) HET-1 is pyrazolyl (optionally substituted with ethyl, isopropyl, difluoromethyl, or 1 or 2 methyl), thiazolyl (optionally substituted with methyl), pyrazinyl (optionally substituted with methyl), pyridyl (optionally substituted by fluoro), isoxazolyl (optionally substituted with methyl) and thiadiazolyl (optionally substituted with methyl)

- 12 -

(43) HET-1 is selected from pyrazinyl (optionally substituted with methyl), pyrazolyl (optionally substituted on carbon by methyl), methylthiadiazolyl (particularly 1,2,4-thiadiazol-5-yl, more particularly 3-methyl-1,2,4-thiadiazol-5-yl), thiazolyl (optionally substituted with methyl), pyridyl (optionally substituted by fluoro) and isoxazolyl

5 (44) R⁶ is selected from (1-4C)alkyl, halo, hydroxy(1-4C)alkyl, di(1-4C)alkylamino(1-4C)alkyl

(45) R⁶ is selected from methyl, ethyl, chloro, fluoro, hydroxymethyl, methoxymethyl, aminomethyl, N-methylaminomethyl, dimethylaminomethyl

(46) R⁶ is selected from methyl, ethyl, chloro, fluoro, hydroxymethyl and methoxymethyl

10 (47) R⁶ is selected from methyl or ethyl

(48) R⁶ is methyl

(49) R⁶ is selected from (1-4C)alkyl and (1-4C)alkoxy(1-4C)alkyl

(50) R⁶ is selected from methyl, ethyl, isopropyl and methoxymethyl

15 (51) when 2 substituents R⁶ are present, both are selected from methyl, ethyl, bromo, chloro and fluoro; preferably both are methyl and at least one is on an available nitrogen atom

(52) R⁴ is hydrogen

20 (53) R⁴ is (1-4C)alkyl [substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and -C(O)NR⁵R⁵]

(54) R⁴ is (1-4C)alkyl [substituted by 1 substituent selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl and -C(O)NR⁵R⁵]

(55) R⁴ is (1-4C)alkyl

(56) R⁴ is (1-4C)alkyl substituted by -OR⁵

25 (57) R⁴ is (1-4C)alkyl substituted by HET-2

(58) R⁴ is (3-6C)cycloalkyl, particularly cyclopropyl or cyclobutyl

(59) R⁴ is (3-6C)cycloalkyl substituted by a group selected from R⁷

(60) R⁴ is (3-6C)cycloalkyl substituted by a group selected from -OR⁵ and (1-4C)alkyl

(61) R⁴ is selected from (1-4C)alkyl and (3-6C)cycloalkyl

30 (62) R⁴ is selected from methyl, ethyl, cyclopropyl and cyclobutyl

(63) R⁴ is HET-2

(64) R⁴ is selected from hydrogen, (1-4C)alkyl, and (1-4C)alkyl substituted with -OR⁵

- 13 -

(65) HET-2 is unsubstituted

(66) HET-2 is substituted with 1 or 2 substituents independently selected from (1-4C)alkyl, hydroxy and (1-4C)alkoxy

(67) HET-2 is a fully saturated ring system

5 (68) HET-2 is a fully unsaturated ring system

(69) HET-2 is selected from azetidinyl, morpholino, morpholinyl, piperidinyl, piperazinyl, 3-oxopiperazinyl, thiomorpholiny, pyrrolidinyl, pyrrolidonyl, 2,5-dioxopyrrolidinyl, 1,1-dioxotetrahydrothienyl, 2-oxazolidinonyl, 2-oxotetrahydrofuranyl, tetrahydrofuranyl, tetrahydropyranyl, 1,1-dioxothiomorpholino, 1,3-dioxolanyl, 2-oxoimidazolidinyl, 2,4-dioxoimidazolidinyl, pyranyl and 4-pyridonyl

(70) HET-2 is selected from azetidinyl, morpholino, morpholinyl, piperidinyl, piperazinyl, pyrrolidinyl, thiomorpholiny, tetrahydrofuranyl, and tetrahydropyranyl

(71) HET-2 is selected from furyl, thienyl, thiazoly, isothiazoly, thiadiazoly, pyridyl, pyrazinyl, pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, 15 oxadiazolyl, pyrrolyl, 1,2,4-triazolyl and 1,2,3-triazolyl

(72) HET-2 is selected from furyl, thienyl, thiazoly, isothiazoly, thiadiazoly, pyridyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl, piperidinyl, piperazinyl, 3-oxopiperazinyl, pyrrolidinyl, pyrrolidonyl, 2-oxazolidinonyl, tetrahydrofuranyl, tetrahydropyranyl, 1,1-dioxotetrahydrothienyl, and 2-oxoimidazolidinyl

20 (73) HET-2 is selected from morpholino, furyl, imidazolyl, oxazolyl, isoxazolyl, oxadiazolyl, piperidinyl, piperazinyl, 3-oxopiperazinyl, pyrrolidinyl, 2-pyrrolidonyl, 2-oxazolidinonyl, tetrahydrofuranyl, tetrahydropyranyl, 1,1-dioxotetrahydrothienyl, and 2-oxoimidazolidinyl

(74) HET-2 is selected from morpholino, furyl, imidazolyl, isoxazolyl, oxadiazolyl, 25 piperidinyl, piperazinyl, 3-oxopiperazinyl, pyrrolidinyl, 2-pyrrolidonyl, tetrahydropyranyl, 1,1-dioxotetrahydrothienyl, and 2-oxoimidazolidinyl

(75) HET-2 is oxadiazolyl or pyrazolyl

(76) R⁵ is hydrogen

(77) R⁵ is (1-4)alkyl, preferably methyl

30 (78) R⁵ is hydrogen or methyl

(79) R⁷ is a substituent on carbon and is selected from hydroxy, (1-4C)alkoxy, (1-4C)alkyl, -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, (1-4C)alkoxy(1-4C)alkyl, and hydroxy(1-4C)alkyl

- 14 -

(80) R⁷ is a substituent on carbon and is selected from hydroxy, (1-4C)alkoxy, (1-4C)alkyl, -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, and hydroxy(1-4C)alkyl

(81) R⁷ is a substituent on carbon and is selected from hydroxy, methoxy, -COMe, -CONH₂, -CONHMe, -CONMe₂, and hydroxymethyl

5 (82) R⁷ is a substituent on carbon and is selected from (1-4C)alkyl, hydroxy and (1-4C)alkoxy

(83) R⁷ is a substituent on carbon and is selected from methyl, ethyl, methoxy and hydroxy

(84) R⁷ is a substituent on nitrogen and is selected from (1-4C)alkyl, -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, (1-4C)alkoxy(1-4C)alkyl, and hydroxy(1-4C)alkyl

10 (85) R⁷ is a substituent on nitrogen and is selected from (1-4C)alkyl, hydroxy and (1-4C)alkoxy

(86) R⁷ is methyl

(87) R⁸ is selected from methyl, hydroxy, methoxy, -CONH₂, -CONHMe, -CONMe₂, hydroxymethyl, hydroxyethyl, -NHMe and -NMe₂

15 (88) R⁸ is selected from methyl, -CONH₂, hydroxyethyl and hydroxy

(89) R⁸ is selected from (1-4C)alkyl and (1-4C)alkoxy

(90) R⁸ is selected from methyl, methoxy and isopropoxy

(91) R⁸ is methyl

(92) R⁹ is selected from methyl, hydroxy, methoxy, -CONH₂, -CONHMe, -CONMe₂, hydroxymethyl, hydroxyethyl, -NHMe and -NMe₂

20 (93) R⁹ is methyl

(94) HET-2 is a 5-membered ring

(95) HET-2 is a 6-membered ring

(96) HET-2 is selected from thienyl, furyl, thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl, pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl and oxadiazolyl

25 (97) HET-2 is selected from thienyl, furyl, thiadiazolyl, pyridyl, pyrazinyl, pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl and oxadiazolyl

(98) HET-2 is selected from pyridyl, pyrazinyl, thiazolyl and thienyl

(99) HET-2 is selected from pyridyl, pyrazinyl and thiazolyl

30 (101) HET-2 is selected from pyridyl, pyrazinyl, pyridazinyl and thiazolyl

(102) HET-2 is selected from pyridyl and pyrazinyl

(103) HET-2 is pyrazinyl

- 15 -

- (104) HET-2 is not substituted on nitrogen
- (105) HET-2 has one nitrogen substituent selected from R⁸
- (106) R⁸ is (1-4C)alkyl
- (107) R⁸ is (3-6C)cycloalkyl
- 5 (108) R⁸ is hydroxy(1-4C)alkyl or (1-4C)alkoxy(1-4C)alkyl
- (109) R⁸ is -C(O)(1-4C)alkyl
- (110) R⁸ is benzyl
- (111) R⁸ is (1-4C)alkylsulfonyl
- (112) R⁸ is (1-4C)alkyl or benzyl

10 According to a further feature of the invention there is provided the following preferred groups of compounds of the invention:

In one aspect of the invention there is provided a compound of formula (I), or a salt thereof, wherein

15 R¹ is selected from fluoromethoxymethyl and difluoromethoxymethyl (particularly difluoromethoxymethyl);

R² is selected from -C(O)NR⁴R⁵, -SO₂NR⁴R⁵ and -SO_pR⁴;

Ring A is pyridyl or pyrazinyl;

R³ is selected from fluoro and chloro;

n is 0 or 1;

20 HET-1 is selected from pyrazolyl (optionally substituted with ethyl, isopropyl, difluoromethyl, or 1 or 2 methyl), thiazolyl (optionally substituted with methyl), pyrazinyl (optionally substituted with methyl), pyridyl (optionally substituted by fluoro), isoxazolyl (optionally substituted with methyl) and thiadiazolyl (optionally substituted with methyl);

R⁴ is hydrogen or (1-4C)alkyl;

25 R⁵ is hydrogen or (1-4C)alkyl;

p is 0, 1 or 2, particularly 2.

In another aspect of the invention there is provided a compound of formula (I), or a salt thereof, wherein

30 R¹ is selected from fluoromethoxymethyl and difluoromethoxymethyl (particularly difluoromethoxymethyl);

R² is selected from -C(O)NR⁴R⁵ and -SO_pR⁴;

Ring A is pyridyl or pyrazinyl;

- 16 -

R³ is selected from fluoro and chloro;

n is 0 or 1;

HET-1 is selected from pyrazolyl (optionally substituted with methyl) and pyrazinyl (optionally substituted with methyl);

5 R⁴ is (1-4C)alkyl;

R⁵ is hydrogen or (1-4C)alkyl;

p is 0, 1 or 2, particularly 2.

In another aspect of the invention there is provided a compound of formula (I), or a salt thereof, wherein

10 R¹ is selected from fluoromethoxymethyl and difluoromethoxymethyl (particularly difluoromethoxymethyl);

R² is selected from -C(O)NR⁴R⁵ and -SO_pR⁴;

Ring A is pyridyl or pyrazinyl;

R³ is selected from fluoro and chloro;

15 n is 0 or 1;

HET-1 is selected from pyrazolyl (optionally substituted with methyl) and pyrazinyl (optionally substituted with methyl);

R⁴ is methyl;

R⁵ is hydrogen or methyl;

20 p is 0, 1 or 2, particularly 2.

Further preferred compounds of the invention are each of the Examples, each of which provides a further independent aspect of the invention. In further aspects, the present invention also comprises any two or more compounds of the Examples.

25 Particular compounds of the invention include any one or more of:

3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-N-(5-methylpyrazin-2-yl)-5-(6-methylsulfonylpyridin-3-yl)oxy-benzamide;

5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide;

5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-(1H-pyrazol-3-ylcarbamoyl)phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide;
3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-(6-methylsulfonylpyridin-3-yl)oxy-N-(1H-pyrazol-3-yl)benzamide;
5 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(1-methylpyrazol-3-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide;
3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-N-(1-methylpyrazol-3-yl)-5-(6-methylsulfonylpyridin-3-yl)oxy-benzamide;
or a salt thereof.

10 The compounds of the invention may be administered in the form of a pro-drug.

A pro-drug is a bioprecursor or pharmaceutically acceptable compound being degradable in the body to produce a compound of the invention (such as an ester or amide of a compound of the invention, particularly an in-vivo hydrolysable ester).

Various forms of prodrugs are known in the art. For examples of such prodrug derivatives, see:

- a) Design of Prodrugs, edited by H. Bundgaard, (Elsevier, 1985) and Methods in Enzymology, Vol. 42, p. 309-396, edited by K. Widder, *et al.* (Academic Press, 1985);
- b) A Textbook of Drug Design and Development, edited by Krosgaard-Larsen;
- c) H. Bundgaard, Chapter 5 "Design and Application of Prodrugs", by H. Bundgaard p. 113-191 (1991);
- d) H. Bundgaard, Advanced Drug Delivery Reviews, 8, 1-38 (1992);
- e) H. Bundgaard, *et al.*, Journal of Pharmaceutical Sciences, 77, 285 (1988); and
- f) N. Kakeya, *et al.*, Chem Pharm Bull, 32, 692 (1984).

The contents of the above cited documents are incorporated herein by reference.

25 Examples of pro-drugs are as follows. An in-vivo hydrolysable ester of a compound of the invention containing a carboxy or a hydroxy group is, for example, a pharmaceutically-acceptable ester which is hydrolysed in the human or animal body to produce the parent acid or alcohol. Suitable pharmaceutically-acceptable esters for carboxy include C₁ to C₆alkoxymethyl esters for example methoxymethyl, C₁ to C₆alkanoyloxymethyl esters for example pivaloyloxymethyl, phthalidyl esters, C₃ to C₈cycloalkoxycarbonyloxyC₁ to C₆alkyl esters for example

1-cyclohexylcarbonyloxyethyl; 1,3-dioxolen-2-onylmethyl esters, for example 5-methyl-1,3-dioxolen-2-onylmethyl; and C₁₋₆alkoxycarbonyloxyethyl esters.

An in-vivo hydrolysable ester of a compound of the invention containing a hydroxy group includes inorganic esters such as phosphate esters (including phosphoramidic cyclic esters) and α -acyloxyalkyl ethers and related compounds which as a result of the in-vivo hydrolysis of the ester breakdown to give the parent hydroxy group/s. Examples of α -acyloxyalkyl ethers include acetoxymethoxy and 2,2-dimethylpropionyloxy-methoxy. A selection of in-vivo hydrolysable ester forming groups for hydroxy include alkanoyl, benzoyl, phenylacetyl and substituted benzoyl and phenylacetyl, alkoxycarbonyl (to give alkyl carbonate esters), dialkylcarbamoyl and N-(dialkylaminoethyl)-N-alkylcarbamoyl (to give carbamates), dialkylaminoacetyl and carboxyacetyl.

Under certain conditions, compounds of Formula (I) may form pharmaceutically acceptable salts. A suitable pharmaceutically-acceptable salt of a compound of the invention is, for example, an acid-addition salt of a compound of the invention which is sufficiently basic, for example, an acid-addition salt with, for example, an inorganic or organic acid, for example hydrochloric, hydrobromic, sulphuric, phosphoric, trifluoroacetic, citric or maleic acid. It will be understood that an acid addition salt may be formed with any sufficiently basic group which may for example be in HET-1 or may for example be a substituent R². In addition a suitable pharmaceutically acceptable salt of a compound of the invention which is sufficiently acidic is an alkali metal salt, for example a sodium or potassium salt, an alkaline earth metal salt, for example a calcium or magnesium salt, an ammonium salt or a salt with an organic base which affords a physiologically-acceptable cation, for example a salt with methylamine, dimethylamine, trimethylamine, piperidine, morpholine or tris-(2-hydroxyethyl)amine.

A further feature of the invention is a pharmaceutical composition comprising a compound of Formula (I) as defined above, or a pharmaceutically-acceptable salt thereof, together with a pharmaceutically-acceptable diluent or carrier.

According to another aspect of the invention there is provided a compound of Formula (I) as defined above or a pharmaceutically-acceptable salt thereof for use as a medicament.

According to another aspect of the invention there is provided a compound of Formula (I), or a pharmaceutically-acceptable salt thereof as defined above for use as a

medicament for treatment of a disease mediated through GLK, in particular type 2 diabetes.

Further according to the invention there is provided the use of a compound of Formula (I) or a pharmaceutically-acceptable salt thereof in the preparation of a medicament for treatment of a disease mediated through GLK, in particular type 2 diabetes.

The compound is suitably formulated as a pharmaceutical composition for use in this way.

According to another aspect of the present invention there is provided a method of treating GLK mediated diseases, especially diabetes, by administering an effective amount of a compound of Formula (I) or a pharmaceutically-acceptable salt thereof, to a mammal in need of such treatment.

According to another aspect of the present invention there is provided the use of a compound of Formula (I), or a pharmaceutically-acceptable salt thereof, for treatment of a disease mediated through GLK.

According to another aspect of the present invention there is provided the use of a compound of Formula (I), or a pharmaceutically-acceptable salt thereof, for treatment of type 2 diabetes.

Specific diseases which may be treated by a compound or composition of the invention include: blood glucose lowering in Type 2 Diabetes Mellitus without a serious risk of hypoglycaemia (and potential to treat type 1), dyslipidemia, obesity, insulin resistance, metabolic syndrome X, impaired glucose tolerance.

As discussed above, thus the GLK/GLKRP system can be described as a potential “Diabesity” target (of benefit in both Diabetes and Obesity). Thus, according to another aspect of the invention there is provided the use of a compound of Formula (I) or a pharmaceutically-acceptable salt thereof, in the preparation of a medicament for use in the combined treatment or prevention, particularly treatment, of diabetes and obesity.

According to another aspect of the invention there is provided the use of a compound of Formula (I) or a pharmaceutically-acceptable salt thereof, in the preparation of a medicament for use in the treatment or prevention of obesity.

According to a further aspect of the invention there is provided a method for the combined treatment of obesity and diabetes by administering an effective amount of a

compound of Formula (I) or a pharmaceutically-acceptable salt thereof, to a mammal in need of such treatment.

According to another aspect of the invention there is provided a compound of Formula (I) or a pharmaceutically-acceptable salt thereof as defined above for use as a medicament for treatment or prevention, particularly treatment of obesity.

According to a further aspect of the invention there is provided a method for the treatment of obesity by administering an effective amount of a compound of Formula (I) or a pharmaceutically-acceptable salt thereof, to a mammal in need of such treatment.

Compounds of the invention may be particularly suitable for use as pharmaceuticals, for example because of favourable physical and/or pharmacokinetic properties and/or toxicity profile.

The compositions of the invention may be in a form suitable for oral use (for example as tablets, lozenges, hard or soft capsules, aqueous or oily suspensions, emulsions, dispersible powders or granules, syrups or elixirs), for topical use (for example as creams, ointments, gels, or aqueous or oily solutions or suspensions), for administration by inhalation (for example as a finely divided powder or a liquid aerosol), for administration by insufflation (for example as a finely divided powder) or for parenteral administration (for example as a sterile aqueous or oily solution for intravenous, subcutaneous, intramuscular or intramuscular dosing or as a suppository for rectal dosing). Dosage forms suitable for oral use are preferred.

The compositions of the invention may be obtained by conventional procedures using conventional pharmaceutical excipients, well known in the art. Thus, compositions intended for oral use may contain, for example, one or more colouring, sweetening, flavouring and/or preservative agents.

Suitable pharmaceutically acceptable excipients for a tablet formulation include, for example, inert diluents such as lactose, sodium carbonate, calcium phosphate or calcium carbonate, granulating and disintegrating agents such as corn starch or algenic acid; binding agents such as starch; lubricating agents such as magnesium stearate, stearic acid or talc; preservative agents such as ethyl or propyl *p*-hydroxybenzoate, and anti-oxidants, such as ascorbic acid. Tablet formulations may be uncoated or coated either to modify their disintegration and the subsequent absorption of the active ingredient within the

gastrointestinal tract, or to improve their stability and/or appearance, in either case, using conventional coating agents and procedures well known in the art.

Compositions for oral use may be in the form of hard gelatin capsules in which the active ingredient is mixed with an inert solid diluent, for example, calcium carbonate, 5 calcium phosphate or kaolin, or as soft gelatin capsules in which the active ingredient is mixed with water or an oil such as peanut oil, liquid paraffin, or olive oil.

Aqueous suspensions generally contain the active ingredient in finely powdered form together with one or more suspending agents, such as sodium carboxymethylcellulose, methylcellulose, hydroxypropylmethylcellulose, sodium alginate, 10 polyvinyl-pyrrolidone, gum tragacanth and gum acacia; dispersing or wetting agents such as lecithin or condensation products of an alkylene oxide with fatty acids (for example polyoxethylene stearate), or condensation products of ethylene oxide with long chain aliphatic alcohols, for example heptadecaethyleneoxycetanol, or condensation products of ethylene oxide with partial esters derived from fatty acids and a hexitol such as 15 polyoxyethylene sorbitol monooleate, or condensation products of ethylene oxide with long chain aliphatic alcohols, for example heptadecaethyleneoxycetanol, or condensation products of ethylene oxide with partial esters derived from fatty acids and a hexitol such as polyoxyethylene sorbitol monooleate, or condensation products of ethylene oxide with partial esters derived from fatty acids and hexitol anhydrides, for example polyethylene 20 sorbitan monooleate. The aqueous suspensions may also contain one or more preservatives (such as ethyl or propyl p-hydroxybenzoate, anti-oxidants (such as ascorbic acid), colouring agents, flavouring agents, and/or sweetening agents (such as sucrose, saccharine or aspartame).

Oily suspensions may be formulated by suspending the active ingredient in a 25 vegetable oil (such as arachis oil, olive oil, sesame oil or coconut oil) or in a mineral oil (such as liquid paraffin). The oily suspensions may also contain a thickening agent such as beeswax, hard paraffin or cetyl alcohol. Sweetening agents such as those set out above, and flavouring agents may be added to provide a palatable oral preparation. These compositions may be preserved by the addition of an anti-oxidant such as ascorbic acid.

30 Dispersible powders and granules suitable for preparation of an aqueous suspension by the addition of water generally contain the active ingredient together with a dispersing or wetting agent, suspending agent and one or more preservatives. Suitable dispersing or

wetting agents and suspending agents are exemplified by those already mentioned above. Additional excipients such as sweetening, flavouring and colouring agents, may also be present.

The pharmaceutical compositions of the invention may also be in the form of oil-in-water emulsions. The oily phase may be a vegetable oil, such as olive oil or arachis oil, or a mineral oil, such as for example liquid paraffin or a mixture of any of these. Suitable emulsifying agents may be, for example, naturally-occurring gums such as gum acacia or gum tragacanth, naturally-occurring phosphatides such as soya bean, lecithin, an esters or partial esters derived from fatty acids and hexitol anhydrides (for example sorbitan monooleate) and condensation products of the said partial esters with ethylene oxide such as polyoxyethylene sorbitan monooleate. The emulsions may also contain sweetening, flavouring and preservative agents.

Syrups and elixirs may be formulated with sweetening agents such as glycerol, propylene glycol, sorbitol, aspartame or sucrose, and may also contain a demulcent, preservative, flavouring and/or colouring agent.

The pharmaceutical compositions may also be in the form of a sterile injectable aqueous or oily suspension, which may be formulated according to known procedures using one or more of the appropriate dispersing or wetting agents and suspending agents, which have been mentioned above. A sterile injectable preparation may also be a sterile injectable solution or suspension in a non-toxic parenterally-acceptable diluent or solvent, for example a solution in 1,3-butanediol.

Compositions for administration by inhalation may be in the form of a conventional pressurised aerosol arranged to dispense the active ingredient either as an aerosol containing finely divided solid or liquid droplets. Conventional aerosol propellants such as volatile fluorinated hydrocarbons or hydrocarbons may be used and the aerosol device is conveniently arranged to dispense a metered quantity of active ingredient.

For further information on formulation the reader is referred to Chapter 25.2 in Volume 5 of Comprehensive Medicinal Chemistry (Corwin Hansch; Chairman of Editorial Board), Pergamon Press 1990.

The amount of active ingredient that is combined with one or more excipients to produce a single dosage form will necessarily vary depending upon the host treated and the particular route of administration. For example, a formulation intended for oral

administration to humans will generally contain, for example, from 0.5 mg to 2 g of active agent compounded with an appropriate and convenient amount of excipients which may vary from about 5 to about 98 percent by weight of the total composition. Dosage unit forms will generally contain about 1 mg to about 500 mg of an active ingredient. For 5 further information on Routes of Administration and Dosage Regimes the reader is referred to Chapter 25.3 in Volume 5 of Comprehensive Medicinal Chemistry (Corwin Hansch; Chairman of Editorial Board), Pergamon Press 1990.

The size of the dose for therapeutic or prophylactic purposes of a compound of the Formula (I) will naturally vary according to the nature and severity of the conditions, the 10 age and sex of the animal or patient and the route of administration, according to well known principles of medicine.

In using a compound of the Formula (I) for therapeutic or prophylactic purposes it will generally be administered so that a daily dose in the range, for example, 0.5 mg to 75 mg per kg body weight is received, given if required in divided doses. In general lower 15 doses will be administered when a parenteral route is employed. Thus, for example, for intravenous administration, a dose in the range, for example, 0.5 mg to 30 mg per kg body weight will generally be used. Similarly, for administration by inhalation, a dose in the range, for example, 0.5 mg to 25 mg per kg body weight will be used. Oral administration is however preferred.

20 The elevation of GLK activity described herein may be applied as a sole therapy or in combination with one or more other substances and/or treatments for the indication being treated. Such conjoint treatment may be achieved by way of the simultaneous, sequential or separate administration of the individual components of the treatment. Simultaneous treatment may be in a single tablet or in separate tablets. For example in the 25 treatment of diabetes mellitus, chemotherapy may include the following main categories of treatment:

- 1) Insulin and insulin analogues;
- 2) Insulin secretagogues including sulphonylureas (for example glibenclamide, glipizide), prandial glucose regulators (for example repaglinide, nateglinide);
- 30 3) Agents that improve incretin action (for example dipeptidyl peptidase IV inhibitors, and GLP-1 agonists);

- 4) Insulin sensitising agents including PPARgamma agonists (for example pioglitazone and rosiglitazone), and agents with combined PPARalpha and gamma activity;
- 5) Agents that modulate hepatic glucose balance (for example metformin, fructose 1, 6 bisphosphatase inhibitors, glycogen phosphorylase inhibitors, glycogen synthase kinase inhibitors);
- 6) Agents designed to reduce the absorption of glucose from the intestine (for example acarbose);
- 7) Agents that prevent the reabsorption of glucose by the kidney (SGLT inhibitors);
- 10) Agents designed to treat the complications of prolonged hyperglycaemia (for example aldose reductase inhibitors);
- 9) Anti-obesity agents (for example sibutramine and orlistat);
- 10) Anti- dyslipidaemia agents such as, HMG-CoA reductase inhibitors (eg statins); PPAR α agonists (fibrates, eg gemfibrozil); bile acid sequestrants (cholestyramine);
- 15) cholesterol absorption inhibitors (plant stanols, synthetic inhibitors); bile acid absorption inhibitors (IBATi) and nicotinic acid and analogues (niacin and slow release formulations);
- 11) Antihypertensive agents such as, β blockers (eg atenolol, inderal); ACE inhibitors (eg lisinopril); Calcium antagonists (eg. nifedipine); Angiotensin receptor antagonists (eg candesartan), α antagonists and diuretic agents (eg. furosemide, benzthiazide);
- 20) 12) Haemostasis modulators such as, antithrombotics, activators of fibrinolysis and antiplatelet agents; thrombin antagonists; factor Xa inhibitors; factor VIIa inhibitors); antiplatelet agents (eg. aspirin, clopidogrel); anticoagulants (heparin and Low molecular weight analogues, hirudin) and warfarin;
- 13) Agents which antagonise the actions of glucagon; and
- 25) 14) Anti-inflammatory agents, such as non-steroidal anti-inflammatory drugs (eg. aspirin) and steroid anti-inflammatory agents (eg. cortisone).

According to another aspect of the present invention there is provided individual compounds produced as end products in the Examples set out below and salts thereof.

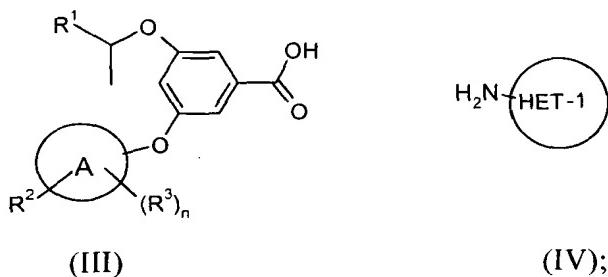
A compound of the invention, or a salt thereof, may be prepared by any process known to be applicable to the preparation of such compounds or structurally related compounds. Functional groups may be protected and deprotected using conventional methods. For examples of protecting groups such as amino and carboxylic acid protecting

- 25 -

groups (as well as means of formation and eventual deprotection), see T.W. Greene and P.G.M. Wuts, "Protective Groups in Organic Synthesis", Second Edition, John Wiley & Sons, New York, 1991.

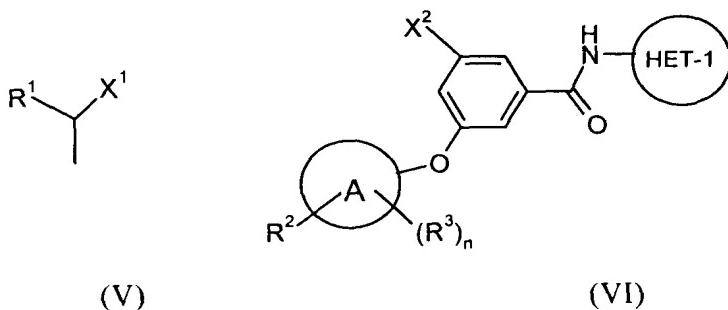
Processes for the synthesis of compounds of Formula (I) are provided as a further feature of the invention. Thus, according to a further aspect of the invention there is provided a process for the preparation of a compound of Formula (I), which comprises a process a) to e) (wherein the variables are as defined hereinbefore for compounds of Formula (I) unless otherwise defined):

(a) reaction of an acid of Formula (III) or activated derivative thereof with a compound of Formula (IV), wherein R¹ is as defined for formula (I) or is a precursor thereof;



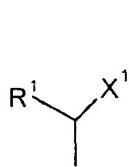
or

(b) reaction of a compound of Formula (V) with a compound of Formula (VI),

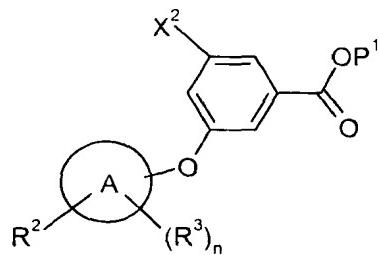


wherein X^1 is a leaving group and X^2 is a hydroxyl group or X^1 is a hydroxyl group and X^2 is a leaving group, and wherein R^1 is as defined for formula (I) or is a precursor thereof; process (b) could also be accomplished using the intermediate ester Formula (VII), wherein P^1 is a protecting group as hereinafter described, followed by ester hydrolysis and amide formation by procedures described elsewhere and well known to those skilled in the art;

- 26 -



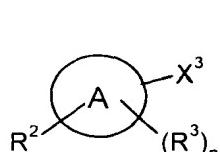
(V)



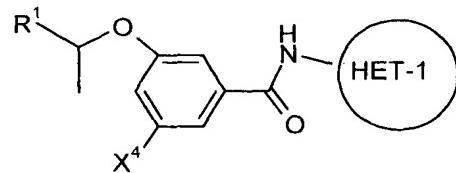
(VII)

or

(c) reaction of a compound of Formula (VIII) with a compound of Formula (IX)



(VIII)

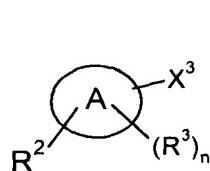


(IX)

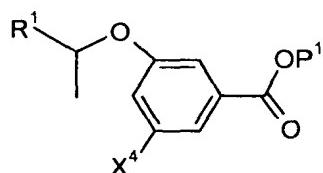
5

wherein X³ is a leaving group or an organometallic reagent and X⁴ is a hydroxyl group or X³ is a hydroxyl group and X⁴ is a leaving group or an organometallic reagent, and wherein R¹ is as defined for formula (I) or is a precursor thereof;

10 process (c) could also be accomplished using the intermediate ester Formula (X), followed by ester hydrolysis and amide formation by procedures described elsewhere and well known to those skilled in the art;



(VIII)

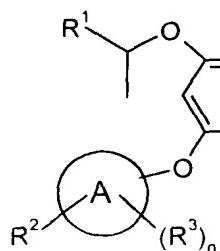


(X)

15

or

(d) reaction of a compound of Formula (XI) with a compound of Formula (XII),



(XI)

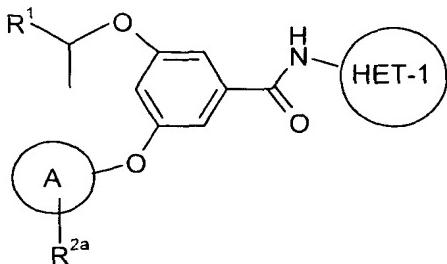


(XII);

- 27 -

wherein X^5 is a leaving group; and wherein R^1 is as defined for formula (I) or is a precursor thereof; or

e) reaction of a compound of formula (XIII)



(XIII)

5 wherein R^{2a} is a precursor to R^2 as $-\text{CONR}^4\text{R}^5$ or $-\text{SO}_2\text{R}^4\text{R}^5$, such as a carboxylic acid, ester or anhydride (for $R^2 = -\text{CONR}^4\text{R}^5$) or the sulfonic acid equivalents (for R^2 is $-\text{SO}_2^2\text{NR}^4\text{R}^5$); with an amine of formula $-\text{NR}^4\text{R}^5$;

and thereafter, if necessary:

- 10 i) converting a compound of Formula (I) into another compound of Formula (I);
- ii) converting a precursor of R^1 into R^1 ;
- iii) removing any protecting groups; and/or
- iv) forming a salt thereof.

Suitable leaving groups X^1 to X^5 for processes b) to d) are any leaving group known in
15 the art for these types of reactions, for example halo, alkoxy, trifluoromethanesulfonyloxy, methanesulfonyloxy, or p-toluenesulfonyloxy; or a group (such as a hydroxy group) that may be converted into a leaving group (such as an oxytriphenylphosphonium group) *in situ*.

Suitable precursors to R^1 include a hydroxy group or a protected hydroxy group,
such as any suitable protected hydroxy group known in the art, for example simple ethers
20 such as a methyl ether, or silyl ethers such as $-\text{OSi}[(1-4\text{C})\text{alkyl}]_3$ (wherein each (1-4C)alkyl group is independently selected from methyl, ethyl, propyl, isopropyl, and tertbutyl).

Examples of such trialkylsilyl groups are trimethylsilyl, triethylsilyl, triisopropylsilyl and
tert-butyldimethylsilyl. Further suitable silyl ethers are those containing phenyl and
substituted phenyl groups, such as $-\text{Si}(\text{PhMe}_2)_2$ and $-\text{Si}(\text{TolMe}_2)_2$ (wherein Tol =
25 methylbenzene). Further suitable values for hydroxy protecting groups are given
hereinafter. R^1 itself may then be generated by removing the hydroxy protecting group if
present, and then by reacting with, for example 2-(fluorosulphonyl)difluoroacetic acid in

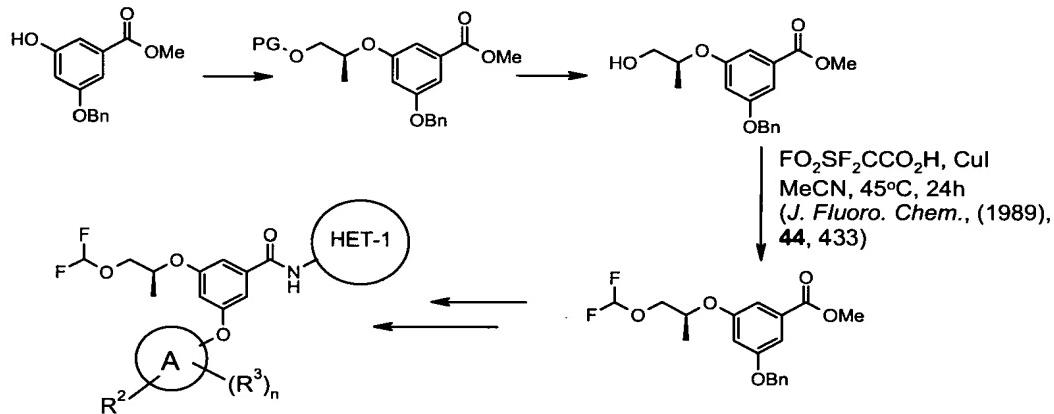
the presence of copper (I)iodide to give the compound wherein R¹ is difluoromethoxymethyl. This reaction is illustrated in Scheme 1. Other values of R¹ may be generated similarly or by methods well known in the art, see for example Bull. Chem. Soc. Japan, 73 (2000), 471-484, 471-484, International Patent application WO 5 2002/050003 and Bioorganic and Medicinal Chemistry Letters, (2001), 11, 407.

Compounds of Formulae (III) to (XII) are commercially available, or are known in the art, or may be made by processes known in the art, for example as shown in the accompanying Examples. For further information on processes for making such compounds, we refer to our PCT publications WO 03/000267, WO 03/015774, WO 03/000262, WO 2004/076420, WO 10 2005/054200, WO 2005/054233, WO 2005/044801 and WO 2005/056530 and references therein. In general it will be appreciated that any aryl-O or alkyl-O bond may be formed by nucleophilic substitution or metal catalysed processes, optionally in the presence of a suitable base.

Compounds of Formula (XIII) may be made by processes such as those shown in processes a) to d) and/or by those processes mentioned above for compounds of formulae (III) to (XII).

Compounds of formulae (III), (IX), (X), (XI) and (XIII) may be made by reaction of suitable precursors with compounds of formula (V) or derivatives thereof, depending on the nature of the R¹ group or its precursor, for example, by nucleophilic displacement of a leaving group X¹ in a compound of formula (V). Compounds of formula (V) are generally commercially available or may be made by simple functional group interconversions from commercially available compounds, or by literature methods. Where the compound of formula (V) contains a precursor to R¹, the R¹ group may be generated in the compound of formula (III), (IX), (X), (XI) or (XIII) as appropriate using reactions such as those illustrated in 20 Schemes 1 and 2 below. Illustrative examples are shown in Schemes 1 and 2 below, and/or in 25 the accompanying examples.

- 29 -

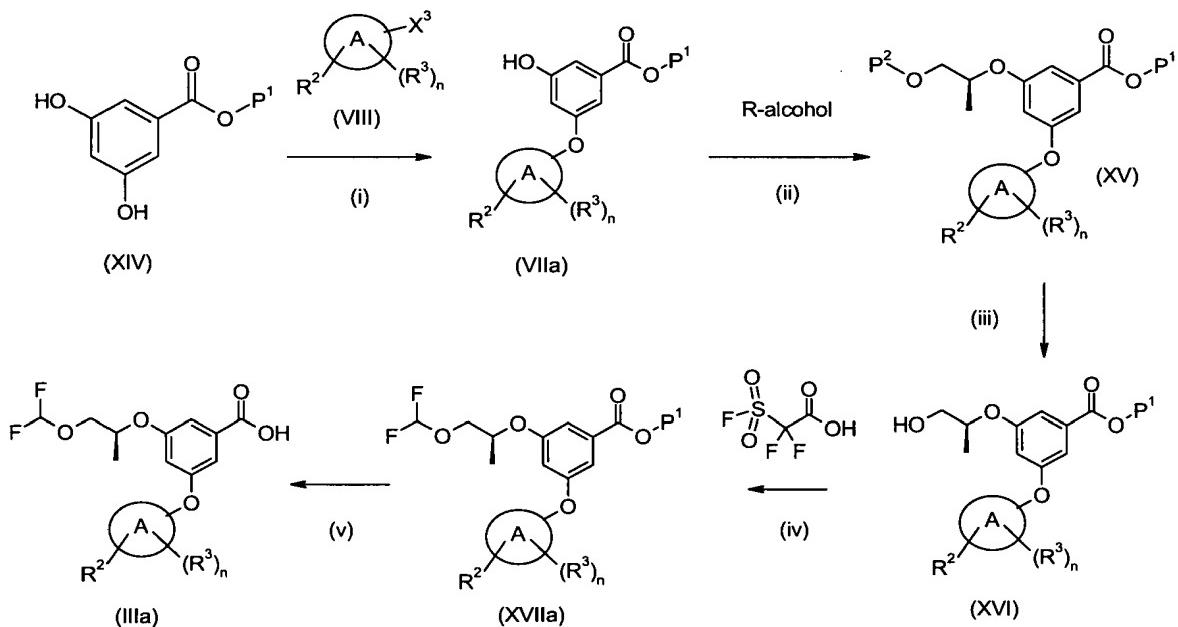


Scheme 1

wherein:

PG is protecting group and R², R³, A, n and HET-1 are as defined for Formula (I).

5



Scheme 2

wherein:

- 10 R², R³, A and n are as defined for Formula (I), P¹ and P² are suitable protecting groups, for example (1-4C)alkyl, and X³ is a leaving group, for example chloro. Suitable reaction conditions for steps (i) to (v) of Scheme 2 are as follows:

- 30 -

Step (i) involves the reaction of Formula (XIV) with a compound of Formula (VIII), for example 5-chloro-N,N-dimethyl-pyrazine-2-carboxamide, in the presence of a suitable base, for example cesium carbonate, in a suitable solvent, for example DMSO, and at a suitable temperature, for example 0 to 60°C, more suitably about 50°C.

5 Step (ii) involves the reaction of a compound of Formula (VII) with an R-alcohol, for example (2R)-1-[(2-methylpropan-2-yl)oxy]propan-2-ol, in the presence of a suitable phosphine, for example triphenylphosphine, and an azodicarboxylate, for example diethylazodicarboxylate, in a suitable solvent, for example THF, and at a suitable temperature, for example 0 to 10°C, more suitably about 0°C.

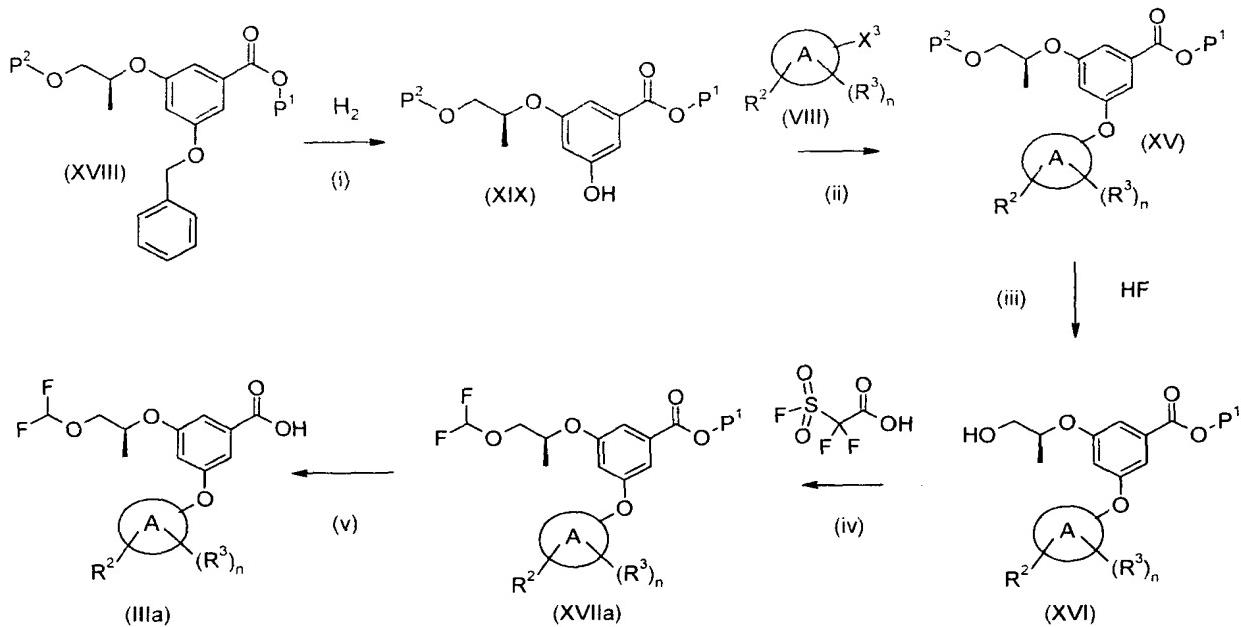
10 Step (iii) involves heating a solution of Formula (XV) in a suitable acid, for example formic acid, at a suitable temperature, for example 0 to 50 to 100°C, more suitably about 90°C.

15 Step (iv) involves the reaction of a compound of Formula (XVI) with 2,2-difluoro-2-fluorosulfonyl-acetic acid, in the presence of a suitable catalyst, for example copper (I) iodide, in a suitable solvent, for example acetonitrile, and at a suitable temperature, for example 0 to 100°C, more suitably about 55°C.

Step (v) involves the reaction of a compound of Formula (XVIIa) with a suitable base, for example NaOH, in a suitable solvent, for example NMP and water, and at a suitable temperature, for example 0 to 25°C, more suitably about 0°C.

20 Alternatively, compounds of Formula (IIIa) where R¹ represents difluoromethoxymethyl may be made according to Scheme 3 as shown below.

- 31 -



wherein:

R^2 , R^3 , A and n are as defined for Formula (I), P^1 and P^2 are suitable protecting groups, for example (1-4C)alkyl and TIPS respectively, and X^3 is a leaving group, for example chloro.

5 Suitable reaction conditions for steps (i) to (v) of Scheme 2 are as follows:

Step (i) involves the reaction of a compound of Formula (XVIII) with hydrogen in the presence of a suitable catalyst, for example 10 % palladium on activated carbon, and at a suitable temperature, for example 0 to 25°C, more suitably about 21°C.

10 Step (ii) involves the reaction of a compound of Formula (XIX) with a compound of Formula (VIII), for example 5-chloro-N,N-dimethyl-pyrazine-2-carboxamide, in the presence of a suitable base, for example potassium carbonate, in a suitable solvent, for example acetonitrile, and at a suitable temperature, for example 0 to 200°C, more suitably about reflux temperature.

15 Step (iii) involves the reaction of a compound of Formula (XV) with hydrogen fluoride in a suitable solvent, for example THF, and at a suitable temperature, for example 0 to 25°C, more suitably about 21°C.

Step (iv) involves the reaction of a compound of Formula (XVI) with 2,2-difluoro-2-fluorosulfonyl-acetic acid, in the presence of a suitable catalyst, for example copper (I) iodide, in a suitable solvent, for example acetonitrile, and at a suitable temperature, for example 0 to 20 100°C, more suitably about 55°C.

Step (v) involves the reaction of a compound of Formula (XVIIa) with a suitable base, for example LiOH, in a suitable solvent, for example THF and methanol, and at a suitable temperature, for example 0 to 25°C, more suitably about 21°C.

Examples of conversions of a compound of Formula (I) into another compound of Formula (I) well known to those skilled in the art, include functional group interconversions such as hydrolysis, hydrogenation, hydrogenolysis, oxidation or reduction, and/or further functionalisation by standard reactions such as amide or metal-catalysed coupling, or nucleophilic displacement reactions. An example would be removal of an R³=chloro substituent, for example by reaction with hydrogen at atmospheric or elevated pressure, in a suitable solvent such as THF/methanol or ethanol.

It will be understood that substituents R², R³ and/or R⁶ may be introduced into the molecule at any convenient point in the synthetic sequence or may be present in the starting materials. A precursor to one of these substituents may be present in the molecule during the process steps a) to e) above, and then be transformed into the desired substituent as a final step to form the compound of formula (I); followed where necessary by

- i) converting a compound of Formula (I) into another compound of Formula (I);
- ii) converting a precursor of R¹ into R¹;
- iii) removing any protecting groups; and/or
- iv) forming a salt thereof.

Specific reaction conditions for the above reactions are as follows, wherein when P¹ is a protecting group P¹ is preferably (1-4C)alkyl, for example methyl or ethyl:

Process a) – coupling reactions of amino groups with carboxylic acids to form an amide are well known in the art. For example,

(i) using an appropriate coupling reaction, such as a carbodiimide coupling reaction performed with EDAC (1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride) in the presence of dimethylaminopyridine (DMAP) in a suitable solvent such as

dichloromethane (DCM), chloroform or dimethylformamide (DMF) at room temperature; or

(ii) reaction in which the carboxylic group is activated to an acid chloride by reaction with oxalyl chloride or 1-chloro-N,N,2-trimethyl-1-propenylamine in the presence of a suitable solvent such as DCM or DMF. The acid chloride can then be reacted with a compound of

Formula (IV) in the presence of a base, such as triethylamine or pyridine, in a suitable solvent such as chloroform or DCM at a temperature between 0°C and 80°C.

Process b) – compounds of Formula (V) and (VI) can be reacted together in a suitable solvent, such as DMF or tetrahydrofuran (THF), with a base such as sodium hydride or potassium *tert*-butoxide, at a temperature in the range 0 to 200°C, optionally using microwave heating or metal catalysis such as palladium(II)acetate, palladium on carbon, copper(II)acetate or copper(I)iodide; alternatively, compounds of Formula (V) and (VI) can be reacted together in a suitable solvent, such as THF or DCM, with a suitable phosphine such as triphenylphosphine, and an azodicarboxylate such as

diethylazodicarboxylate; process b) could also be carried out using a precursor to the ester of formula (VII) such as an aryl-nitrile or trifluoromethyl derivative, followed by conversion to a carboxylic acid and amide formation as previously described;

Process c) - compounds of Formula (VIII) and (IX) can be reacted together in a suitable solvent, such as DMF or THF, with a base such as sodium hydride or potassium *tert*-butoxide, at a temperature in the range 0 to 200°C, optionally using microwave heating or metal catalysis such as palladium(II)acetate, palladium on carbon, copper(II)acetate, copper(I)iodide or bromotris(triphenylphosphine)copper(I); process c) could also be carried out using a precursor to the ester of formula (X) such as an aryl-nitrile or trifluoromethyl derivative, followed by conversion to a carboxylic acid and amide formation as previously described;

Process d) – reaction of a compound of Formula (XI) with a compound of Formula (XII) can be performed in a polar solvent, such as DMF or a non-polar solvent such as THF with a strong base, such as sodium hydride or potassium *tert*-butoxide at a temperature between 0 and 200°C, optionally using microwave heating or metal catalysis, such as

palladium(II)acetate, palladium on carbon, copper(II)acetate or copper(I)iodide;

Process e) - coupling reactions of amino groups with carboxylic or sulfonic acids or acid derivatives to form an amide are well known in the art and are described above for Process a).

Certain intermediates of formula (III), (VI), (VII), (IX) and/or (XI) are believed to be novel and comprise an independent aspect of the invention.

Certain intermediates of formula (III), (IX) and/or (XI) wherein R¹ is as defined herein, are believed to be novel and comprise an independent aspect of the invention.

Certain intermediates of formula (XIII) are believed to be novel and comprise an independent aspect of the invention.

During the preparation process, it may be advantageous to use a protecting group for a functional group within the molecule. Protecting groups may be removed by any convenient method as described in the literature or known to the skilled chemist as appropriate for the removal of the protecting group in question, such methods being chosen so as to effect removal of the protecting group with minimum disturbance of groups elsewhere in the molecule.

Specific examples of protecting groups are given below for the sake of convenience, in which "lower" signifies that the group to which it is applied preferably has 1-4 carbon atoms. It will be understood that these examples are not exhaustive. Where specific examples of methods for the removal of protecting groups are given below these are similarly not exhaustive. The use of protecting groups and methods of deprotection not specifically mentioned is of course within the scope of the invention.

A carboxy protecting group may be the residue of an ester-forming aliphatic or araliphatic alcohol or of an ester-forming silanol (the said alcohol or silanol preferably containing 1-20 carbon atoms). Examples of carboxy protecting groups include straight or branched chain (1-12C)alkyl groups (e.g. isopropyl, *t*-butyl); lower alkoxy lower alkyl groups (e.g. methoxymethyl, ethoxymethyl, isobutoxymethyl; lower aliphatic acyloxy lower alkyl groups, (e.g. acetoxyethyl, propionyloxymethyl, butyryloxymethyl, pivaloyloxymethyl); lower alkoxy carbonyloxy lower alkyl groups (e.g. 1-methoxycarbonyloxyethyl, 1-ethoxycarbonyloxyethyl); aryl lower alkyl groups (e.g. *p*-methoxybenzyl, *o*-nitrobenzyl, *p*-nitrobenzyl, benzhydryl and phthalidyl); tri(lower alkyl)silyl groups (e.g. trimethylsilyl and *t*-butyldimethylsilyl); tri(lower alkyl)silyl lower alkyl groups (e.g. trimethylsilylethyl); and (2-6C)alkenyl groups (e.g. allyl and vinyl ethyl).

Methods particularly appropriate for the removal of carboxyl protecting groups include for example acid-, metal- or enzymically-catalysed hydrolysis.

Examples of hydroxy protecting groups include methyl, *t*-butyl, lower alkenyl groups (e.g. allyl); lower alkanoyl groups (e.g. acetyl); lower alkoxy carbonyl groups (e.g. *t*-butoxycarbonyl); lower alkenyloxy carbonyl groups (e.g. allyloxycarbonyl); aryl lower alkoxy carbonyl groups (e.g. benzoyloxycarbonyl, *p*-methoxybenzyloxycarbonyl, *o*-nitrobenzyloxycarbonyl, *p*-nitrobenzyloxycarbonyl); tri lower alkyl/arylsilyl groups (e.g.

trimethylsilyl, *t*-butyldimethylsilyl, *t*-butyldiphenylsilyl); tetrahydropyran-2-yl; aryl lower alkyl groups (e.g. benzyl) groups; and triaryl lower alkyl groups (e.g. triphenylmethyl).

Examples of amino protecting groups include formyl, aralkyl groups (e.g. benzyl and substituted benzyl, e.g. *p*-methoxybenzyl, nitrobenzyl and 2,4-dimethoxybenzyl, and 5 triphenylmethyl); di-*p*-anisylmethyl and furylmethyl groups; lower alkoxy carbonyl (e.g. *t*-butoxycarbonyl); lower alkenyloxycarbonyl (e.g. allyloxycarbonyl); aryl lower alkoxy carbonyl groups (e.g. benzyloxycarbonyl, *p*-methoxybenzyloxycarbonyl, *o*-nitrobenzyloxycarbonyl, *p*-nitrobenzyloxycarbonyl; trialkylsilyl (e.g. trimethylsilyl and *t*-butyldimethylsilyl); alkylidene (e.g. methylidene); benzylidene and substituted 10 benzylidene groups.

Methods appropriate for removal of hydroxy and amino protecting groups include, for example, nucleophilic displacement, acid-, base, metal- or enzymically-catalysed hydrolysis, catalytic hydrogenolysis/hydrogenation or photolytically for groups such as *o*-nitrobenzyloxycarbonyl, or with fluoride ions for silyl groups. For example, methylether 15 protecting groups for hydroxy groups may be removed by trimethylsilyliodide. A tert-butyl ether protecting group for a hydroxy group may be removed by hydrolysis, for example by use of hydrochloric acid in methanol.

Examples of protecting groups for amide groups include aralkoxymethyl (e.g. benzyloxymethyl and substituted benzyloxymethyl); alkoxy methyl (e.g. methoxymethyl 20 and trimethylsilylethoxymethyl); tri alkyl/arylsilyl (e.g. trimethylsilyl, *t*-butyldimethylsilyl, *t*-butyldiphenylsilyl); tri alkyl/arylsilyloxy methyl (e.g. *t*-butyldimethylsilyloxy methyl, *t*-butyldiphenylsilyloxy methyl); 4-alkoxyphenyl (e.g. 4-methoxyphenyl); 2,4-di(alkoxy)phenyl (e.g. 2,4-dimethoxyphenyl); 4-alkoxybenzyl (e.g. 4-methoxybenzyl); 25 2,4-di(alkoxy)benzyl (e.g. 2,4-di(methoxy)benzyl); and alk-1-enyl (e.g. allyl, but-1-enyl and substituted vinyl e.g. 2-phenylvinyl).

Aralkoxymethyl, groups may be introduced onto the amide group by reacting the latter group with the appropriate aralkoxymethyl chloride, and removed by catalytic hydrogenation. Alkoxy methyl, tri alkyl/arylsilyl and tri alkyl/silyloxy methyl groups may be introduced by reacting the amide with the appropriate chloride and removing with acid; 30 or in the case of the silyl containing groups, fluoride ions. The alkoxyphenyl and alkoxybenzyl groups are conveniently introduced by arylation or alkylation with an appropriate halide and removed by oxidation with ceric ammonium nitrate. Finally alk-1-

enyl groups may be introduced by reacting the amide with the appropriate aldehyde and removed with acid.

In the above other pharmaceutical composition, process, method, use and medicament manufacture features, the alternative and preferred aspects and embodiments of the compounds of the invention described herein also apply.

The following examples are for illustration purposes and are not intended to limit the scope of this application. Each exemplified compound represents a particular and independent aspect of the invention. In the following non-limiting Examples, unless otherwise stated:

- (i) evaporation were carried out by rotary evaporation under reduced pressure and work-up procedures were carried out after removal of residual solids such as drying agents by filtration;
- (ii) operations were carried out at room temperature, that is in the range 18-25°C and under an atmosphere of an inert gas such as argon or nitrogen;
- (iii) yields are given for illustration only and are not necessarily the maximum attainable;
- (iv) the structures of the end-products of the Formula (I) were confirmed by nuclear (generally proton) magnetic resonance (NMR) and mass spectral techniques; proton magnetic resonance chemical shift values were measured on the delta scale and peak multiplicities are shown as follows: s, singlet; d, doublet; t, triplet; m, multiplet; br, broad; q, quartet; quin, quintet; sextet
- (v) intermediates were not generally fully characterised and purity was assessed by thin layer chromatography (TLC), high-performance liquid chromatography (HPLC), infra-red (IR) or NMR analysis;
- (vi) flash chromatography was carried out on silica unless otherwise stated.

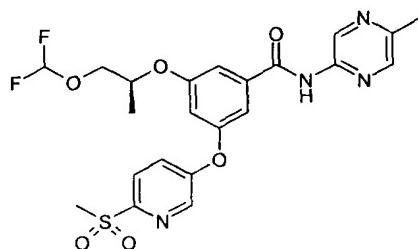
Abbreviations

DCM	dichloromethane;
DEAD	diethylazodicarboxylate;
DIAD	diisopropylazodicarboxylate;
DIPEA	N,N-diisopropylethylamine;
DMA	dimethylacetamide;

- 37 -

DMSO	dimethyl sulfoxide;
DMF	dimethylformamide;
EDAC	1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride;
5 HATU	O-(7-azabenzotriazol-1-yl)-N,N,N',N'-tetramethyluronium hexafluorophosphate;
HPLC	high pressure liquid chromatography;
HPMC	hydroxypropylmethylcellulose;
LCMS	liquid chromatography / mass spectroscopy;
10 NMP	N-methyl-2-pyrrolidone;
NMR	nuclear magnetic resonance spectroscopy;
RT	room temperature;
THF	tetrahydrofuran;
TFA	trifluoroacetic acid;
15 CDCl ₃	deuterochloroform;
MgSO ₄	magnesium sulfate;
PTFE	polytetrafluoroethylene;
TIPS	triisopropylsilyl.

20 **Example 1: 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-N-(5-methylpyrazin-2-yl)-5-(6-methylsulfonylpyridin-3-yl)oxy-benzamide**



A mixture of 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-hydroxy-N-(5-methylpyrazin-2-yl)benzamide (190 mg, 0.54 mmol), 5-bromo-2-methylsulfonyl-pyridine (CAS no. 98626-95-0) (140 mg, 0.59 mmol), cesium carbonate (350 mg, 1.08 mmol) and bromotris(triphenylphosphine)copper(I) (101 mg, 0.11 mmol) in DMA (5 mL) was stirred in a microwave reactor at 160°C for 6 hours. Ethyl acetate (50 mL) was added and the mixture washed with water (50 mL), brine (50 mL), dried (MgSO₄) and reduced *in vacuo*.

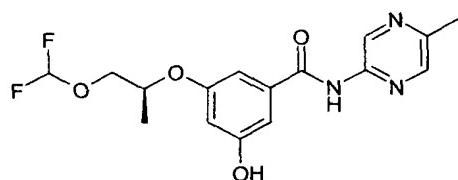
- 38 -

The crude residue was chromatographed on silica, eluting with 10-100% ethyl acetate in isohexane, to give the desired compound (19 mg).

¹H NMR δ (CDCl₃): 1.40 (d, 3H), 2.56 (s, 3H), 3.23 (s, 3H), 3.96 - 4.05 (m, 2H), 4.65 - 4.72 (m, 1H), 6.26 (t, 1H), 6.86 (t, 1H), 7.21 (t, 1H), 7.36 (t, 1H), 7.46 - 7.49 (m, 1H), 8.08 (d, 1H), 8.15 (s, 1H), 8.30 (s, 1H), 8.49 (d, 1H), 9.52 (d, 1H); *m/z* 509 (M+H)⁺

The preparation of 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-hydroxy-N-(5-methylpyrazin-2-yl)benzamide is described below.

¹⁰ 3-[(2S)-1-(Difluoromethoxy)propan-2-yl]oxy-5-hydroxy-N-(5-methylpyrazin-2-yl)benzamide

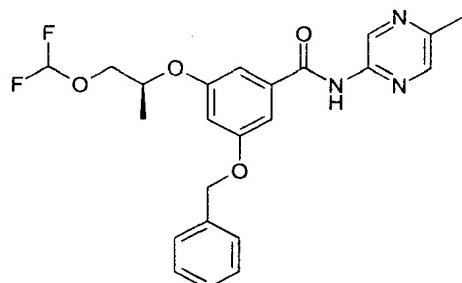


¹⁵ 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-N-(5-methylpyrazin-2-yl)-5-phenylmethoxybenzamide (0.48 g, 1.08 mmol) was dissolved in ethanol (10 mL) and THF (10 mL) and the flask evacuated and purged with argon (3 times). 10% Palladium on carbon (48 mg) was added and the flask further evacuated and finally purged with hydrogen gas. The reaction mixture was stirred at RT for 20 hours. The reaction mixture was evacuated and purged with argon (3 times) and the catalyst removed by filtration through celite®. The filtrate was concentrated *in vacuo* to give the desired compound (0.38 g).

²⁰ ¹H NMR δ (d₆-DMSO): 1.19 (d, 3H), 2.39 (s, 3H), 3.85 - 3.95 (m, 2H), 4.65 - 4.72 (m, 1H), 6.46 (s, 1H), 6.65 (t, 1H), 6.93 (s, 1H), 7.06 (s, 1H), 8.27 (s, 1H), 9.16 (s, 1H), 9.74 (s, 1H), 10.82 (s, 1H); *m/z* 354 (M+H)⁺

- 39 -

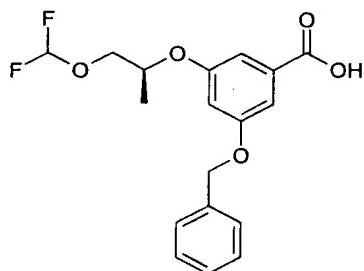
3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-N-(5-methylpyrazin-2-yl)-5-phenylmethoxybenzamide



1-Chloro-N,N,2-trimethyl-prop-1-en-1-amine (0.26 mL, 2.0 mmol) was added to a solution of 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-phenylmethoxy-benzoic acid (0.54 g, 1.5 mmol) in DCM (20 mL) and stirred for 1 hour. 5-Methylpyrazin-2-amine (CAS no. 5521-58-4) (335 mg, 3.1 mmol) then pyridine (0.25 mL, 3.1 mmol) were added and the reaction stirred for a further 30 minutes before being reduced *in vacuo* and partitioned between ethyl acetate (50 mL) and water (50 mL). The aqueous layer was further extracted into ethyl acetate (50 mL) and the combined organics washed with water (50 mL), brine (50 mL), dried (MgSO_4), and reduced *in vacuo*. The crude residue was chromatographed on silica, eluting with 40-100% ethyl acetate in isohexane, to give the desired compound (0.48 g).

^1H NMR δ (CDCl_3): 1.39 (d, 3H), 2.58 (s, 3H), 3.96 - 4.05 (m, 2H), 4.63 - 4.70 (m, 1H), 5.13 (s, 2H), 6.30 (t, 1H), 6.78 (t, 1H), 7.09 (t, 1H), 7.16 (t, 1H), 7.35 - 7.48 (m, 5H), 8.17 (s, 1H), 8.39 (s, 1H), 9.58 (d, 1H); m/z 444 ($\text{M}+\text{H}^+$)⁺

3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-phenylmethoxy-benzoic acid



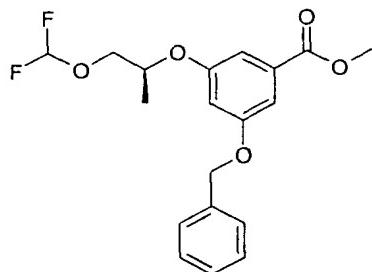
Lithium hydroxide monohydrate (19 mg, 0.45 mmol) in water (2 mL) was added to methyl 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-phenylmethoxy-benzoate (0.11 g, 0.3 mmol) in THF (4 mL) and the mixture stirred at RT for 20 hours. The THF was removed

- 40 -

in vacuo and the aqueous layer adjusted to pH 3 with citric acid then extracted into ethyl acetate (2×30 mL). The organics were washed with water (30 mL), brine (30 mL), dried (MgSO_4), filtered and the solvent removed *in vacuo* to give the desired compound (0.1 g).

$^1\text{H NMR}$ δ (d_6 -DMSO): 1.27 (d, 3H), 4.00 (m, 2H), 4.75 (sextet, 1H), 5.15 (s, 2H), 6.72 (t, 1H), 6.87 (t, 1H), 7.08 (t, 1H), 7.16 (t, 1H), 7.41 (m, 5H), 12.95 (s, 1H); m/z 351 ($M+\text{H}^+$)⁺

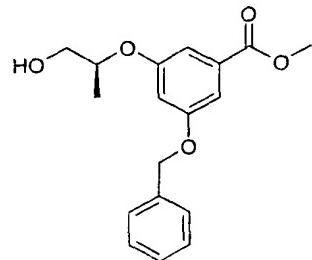
Methyl 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-phenylmethoxy-benzoate



2,2-Difluoro-2-fluorosulfonyl-acetic acid (CAS no. 1717-59-5) (0.239 mL, 2.31 mmol) was added dropwise, with stirring, to a degassed mixture of methyl 3-[(2S)-1-hydroxypropan-2-yl]oxy-5-phenylmethoxy-benzoate (0.73 g, 2.31 mmol) and copper (I) iodide (88 mg, 0.46 mmol) in acetonitrile (10 mL) at 45°C. The reaction was stirred at 45°C for 24 hours. The solvent was removed *in vacuo* and ethyl acetate (30 mL) added. The organics were washed with water (30 mL), brine (30 mL), dried (MgSO_4), filtered and the solvent removed *in vacuo*. The residue was chromatographed on silica, eluting with a gradient of 0-30% ethyl acetate in isohexane, to give the desired compound (0.11 g).

$^1\text{H NMR}$ δ (CDCl_3): 1.37 (d, 3H), 3.93 (s, 3H), 4.00 (m, 2H), 4.63 (sextet, 1H), 5.10 (s, 2H), 6.28 (t, 1H), 6.77 (t, 1H), 7.28 (t, 1H), 7.41 (m, 6H); m/z 367 ($M+\text{H}^+$)⁺

Methyl 3-[(2S)-1-hydroxypropan-2-yl]oxy-5-phenylmethoxy-benzoate



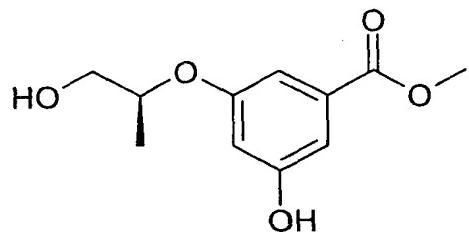
Bromomethylbenzene (1.89 g, 7.20 mmol) was added to a mixture of methyl 3-hydroxy-5-[(2S)-1-hydroxypropan-2-yl]oxy-benzoate (1.55 g, 6.86 mmol) and potassium carbonate

- 41 -

(1.89 g, 0.014 mol) in DMF (16 mL) and the reaction stirred at RT for 20 hours. Ethyl acetate (40 mL) was added and the mixture washed with water (40 mL), saturated sodium bicarbonate solution (40 mL), brine (40 mL), dried (MgSO_4), filtered and the solvent removed *in vacuo*. The residue was chromatographed on silica, eluting with a gradient of 0-100% ethyl acetate in isohexane, to give the desired compound (1.7 g).

^1H NMR δ (CDCl_3): 1.30 (d, 3H), 1.95 (m, 1H), 3.76 (m, 2H), 3.92 (s, 3H), 4.53 (m, 1H), 5.11 (s, 2H), 6.78 (t, 1H), 7.25 (m, 1H), 7.32 (m, 1H), 7.45 (m, 5H); m/z 317 ($\text{M}+\text{H})^+$

Methyl 3-hydroxy-5-[(2S)-1-hydroxypropan-2-yl]oxy-benzoate



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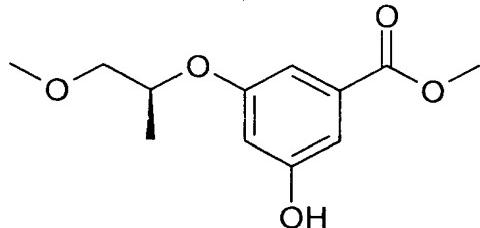
Iodo-trimethyl-silane (CAS no. 16029-98-4) (115 mL, 0.79mol) was added to a solution of methyl 3-hydroxy-5-[(2S)-1-methoxypropan-2-yl]oxy-benzoate (CAS no. 863504-77-2) (38.01 g, 0.158mol) in acetonitrile (500 mL) and stirred for 24 hours. Methanol (300 mL) was added and the reaction stirred for 10 minutes. 10% w/v Aqueous sodium thiosulfate pentahydrate (100 mL) was added to the mixture and stirred for 20 minutes. The reaction mixture was neutralised with saturated aqueous sodium bicarbonate solution, the organic solvents removed *in vacuo*, and the product extracted into ethyl acetate (4×100 mL). The combined organic layers were dried (MgSO_4), filtered and the solvents removed *in vacuo*. The crude material was crystallised from ethyl acetate to give the desired compound (16.8 g).

20

^1H NMR δ (d_6 -DMSO): 1.18 (d, 3H), 3.40-3.55 (m, 2H), 3.80 (s, 3H), 4.35 (sextet, 1H), 4.80 (t, 1H), 6.57 (m, 1H), 6.90 (m, 2H), 9.75 (s, 1H); m/z 304 ($\text{M}+\text{H})^+$

- 42 -

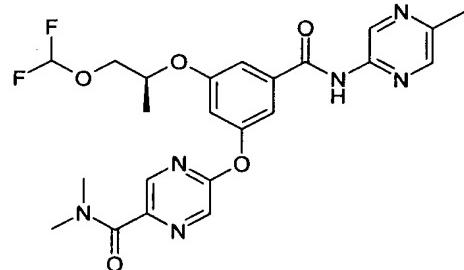
Methyl 3-hydroxy-5-[(2S)-1-methoxypropan-2-yl]oxy-benzoate



Methyl 3-[(2S)-1-methoxypropan-2-yl]oxy-5-phenylimethoxy-benzoate (CAS no. 851885-42-2) (50.0 g, 0.152 mmol) was dissolved in a mixture of THF:ethanol (600 mL) and the flask evacuated and purged with nitrogen (3 times). 10% Palladium on carbon (5.0 g) was added and the flask further evacuated and finally purged with hydrogen gas. The reaction mixture was stirred at ambient temperature for 20 hours until completion. The reaction mixture was evacuated and purged with nitrogen (3 times). The catalyst was filtered off, and the filtrate concentrated *in vacuo* to give the desired compound (36.7 g).

¹⁰ ^1H NMR δ (d₆-DMSO): 1.2 (d, 3H), 3.25 (s, 3H), 3.44 (m, 2H), 3.82 (s, 3H), 4.55 (m, 1H), 6.6 (s, 1H), 6.9 (s, 1H), 6.95 (s, 1H), 9.8 (s, 1H)

Example 2: 5-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide



A mixture of 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-hydroxy-N-(5-methylpyrazin-2-yl)benzamide (0.19 g, 0.54 mmol), 5-chloro-N,N-dimethyl-pyrazine-2-carboxamide (100 mg, 0.54 mmol) and potassium carbonate (149 mg, 1.08 mmol) in acetonitrile (5 mL) was stirred in a microwave reactor at 140°C for 5 hours and the mixture reduced *in vacuo*.

²⁰ Ethyl acetate (50 mL) was added and the mixture washed with water (50 mL), brine (50 mL), dried (MgSO_4), and reduced *in vacuo*. The crude residue was chromatographed on silica, eluting with 0-5% methanol in DCM, to give the desired compound (150 mg).

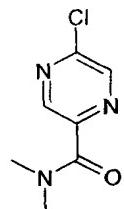
- 43 -

¹H NMR δ (CDCl₃): 1.40 (d, 3H), 2.55 (s, 3H), 3.15 (s, 3H), 3.18 (s, 3H), 3.95 - 4.05 (m, 2H), 4.64 - 4.71 (m, 1H), 6.26 (t, 1H), 6.97 (t, 1H), 7.32 (t, 1H), 7.40 (t, 1H), 8.13 (s, 1H), 8.38 (d, 1H), 8.41 (s, 1H), 8.53 (d, 1H), 9.53 (d, 1H); *m/z* 503 (M+H)⁺

5 The preparation of 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-hydroxy-N-(5-methylpyrazin-2-yl)benzamide was described previously.

The preparation of 5-chloro-N,N-dimethyl-pyrazine-2-carboxamide is described below.

10 5-Chloro-N,N-dimethyl-pyrazine-2-carboxamide



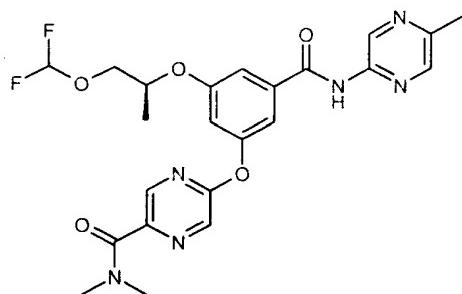
Oxalyl chloride (1.7 mL, 19 mmol) was added to a suspension of 5-chloropyrazine-2-carboxylic acid (CAS no. 36070-80-1) (2.53 g, 16.0 mmol) in dichloromethane (25 mL) and DMF (4 drops) at RT and under argon. The mixture was allowed to stir for 1.5h,

15 concentrated *in vacuo* to and the residue was re-dissolved in dichloromethane (25 mL). Dimethylamine (2M in THF, 8.77 mL, 17.6 mmol) was then added dropwise followed by triethylamine (4.9 mL, 35 mmol) and allowed to stir for a further 5.5 hours. The reaction mixture was concentrated *in vacuo* and the residue was re-dissolved in dichloromethane and filtered. The filtrate was chromatographed on silica, eluting with a gradient of 50-100% ethyl acetate in isohexanes to give the desired compound (1.88 g).

20 ¹H NMR δ (CD₃OD): 3.34 (s, 3H), 3.38 (s, 3H), 8.90 (s, 1H), 8.92 (s, 1H); *m/z* 186 (M+H)⁺

An alternative method for the preparation of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide is given below:

- 44 -



1-Chloro-N,N,2-trimethyl-1-propenylamine (6.9 mL, 52.3 mmol) was added to a solution of 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxy-benzoic acid (17.2 g, 41.8 mmol) in DCM (250 mL) and stirred at ambient temperature for 30 minutes. 5-Methylpyrazin-2-amine (9.1 g, 83.6 mmol) and pyridine (6.8 mL, 83.6 mmol) were added and the reaction stirred overnight. The solvent was evaporated under reduced pressure. The residue was dissolved in ethyl acetate (200 mL), washed with water (2 × 100 mL), citric acid (1N, 100 mL), saturated sodium bicarbonate solution (2 × 100 mL) and saturated brine (10 mL), dried (MgSO_4), filtered and evaporated under reduced pressure. The residue was purified by flash chromatography on silica, eluting with a gradient of 25-100% ethyl acetate in isohexane to afford the product as a colourless foam (11.2 g). To a 200 mg sample of this material was added diethyl ether (1 mL) and the resulting suspension slurried overnight with vigorous stirring. The resulting white solid was isolated by filtration and dried under vacuum. X-ray powder diffraction established this material to contain a large degree of crystalline character. The remaining material (9.2g) was split in to two batches (3.5g and 5.7g). To the smaller batch (3.5g) was added diethyl ether (12.5 mL) and to the larger batch (5.7g) was added diethyl ether (20 mL). The larger batch was seeded with the crystalline material obtained previously (50 mg). Both batches were stirred for 16 hours at room temperature. The resulting colourless solids were isolated by filtration, combined and dried in vacuum. The resulting material (6.1 g, 29%) had an X-ray powder diffraction pattern consistent with the crystalline material obtained previously and with that described for 5-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide Form A described below. ^1H NMR and mass spectrometry data were consistent with those obtained using the previous method.

- 45 -

5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide Form A is characterised in providing at least one of the following 2θ values measured using CuKa radiation: 20.3 and 15.6.

5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide Form A is characterised in providing an X-ray powder diffraction pattern, substantially as shown in Figure A. The ten most prominent peaks are shown in Table A:

Table A

Ten most Prominent X-Ray Powder Diffraction peaks for 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide Form A

Angle 2- Theta (2θ)	Intensity %	Relative Intensity
20.325	100	vs
15.646	94.3	vs
23.15	46.2	vs
22.424	43.8	vs
9.266	39.1	vs
25.707	34.8	vs
26.21	32.9	vs
18.72	28.5	vs
26.485	28.5	vs
8.425	28.5	vs

vs = very strong

According to the present invention there is provided a crystalline form of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide, Form A, which has an X-ray powder diffraction pattern with at least one specific peak at about $2\text{-theta} = 20.3^\circ$.

According to the present invention there is provided a crystalline form, Form A, which has an X-ray powder diffraction pattern with at least one specific peak at about $2\text{-theta} = 15.6^\circ$.

- 46 -

According to the present invention there is provided a crystalline form of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide, Form A, which has an X-ray powder diffraction pattern with at least two specific peaks at about 2-theta = 20.3° and 15.6°.

5 According to the present invention there is provided a crystalline form of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide, Form A, which has an X-ray powder diffraction pattern with specific peaks at about 2-theta = 20.3, 15.6, 23.2, 22.4, 9.3, 25.7, 26.2, 18.7, 26.5 and 8.4°.

10 According to the present invention there is provided crystalline form of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide, Form A, which has an X-ray powder diffraction pattern substantially the same as the X-ray powder diffraction pattern shown in Figure A.

15 According to the present invention there is provided crystalline form of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide, Form A, which has an X-ray powder diffraction pattern with at least one specific peak at 2-theta = 20.3° plus or minus 0.5° 2-theta.

20 According to the present invention there is provided a crystalline form of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide, Form A, which has an X-ray powder diffraction pattern with at least one specific peak at 2-theta = 15.6° plus or minus 0.5° 2-theta.

25 According to the present invention there is provided a crystalline form of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide, Form A, which has an X-ray powder diffraction pattern with at least two specific peaks at 2-theta = 20.3° and 15.6° wherein said values may be plus or minus 0.5° 2-theta.

30 According to the present invention there is provided a crystalline form of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide, Form A, which has an X-ray powder diffraction pattern with specific peaks at 2-theta = 20.3, 15.6, 23.2, 22.4, 9.3, 25.7, 26.2, 18.7, 26.5 and 8.4° wherein said values may be plus or minus 0.5° 2-theta.

According to the present invention there is provided a crystalline form of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide, Form A, which has an X-ray powder diffraction pattern with at least one specific peak at 2-theta = 20.3°.

5 According to the present invention there is provided a crystalline form of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide, Form A, which has an X-ray powder diffraction pattern with at least one specific peak at 2-theta = 15.6°.

10 According to the present invention there is provided a crystalline form of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide, Form A, which has an X-ray powder diffraction pattern with at least two specific peaks at 2-theta = 20.3° and 15.6°.

15 According to the present invention there is provided a crystalline form of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide, Form A, which has an X-ray powder diffraction pattern with specific peaks at 2-theta = 20.3, 15.6, 23.2, 22.4, 9.3, 25.7, 26.2, 18.7, 26.5 and 8.4°.

20 According to the present invention there is provided a crystalline form of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide, Form A, which has an X-ray powder diffraction pattern as shown in Figure A.

25 DSC analysis shows 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide Form A is a low melting solid with an onset of melting at 75.0° C and a peak at 83.1° C (Figure B).

According to the present invention there is therefore provided a crystalline form of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide, Form A, with an onset of melting at about 75.0° C and a peak at about 83.1° C.

30 According to the present invention there is therefore provided a process for the formation of a crystalline form of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide which comprises crystallisation of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-

methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide from a solution of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide in diethyl ether.

When it is stated that the present invention relates to a crystalline form of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide Form A, the degree of crystallinity is conveniently greater than about 60%, more conveniently greater than about 80%, preferably greater than about 90% and more preferably greater than about 95%.

The 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide Form A provides X-ray powder diffraction patterns substantially the same as the X-ray powder diffraction patterns shown in Figure A and has substantially the ten most prominent peaks (angle 2-theta values) shown in Table A. It will be understood that the 2-theta values of the X-ray powder diffraction pattern may vary slightly from one machine to another or from one sample to another, and so the values quoted are not to be construed as absolute.

It is known that an X-ray powder diffraction pattern may be obtained which has one or more measurement errors depending on measurement conditions (such as equipment or machine used). In particular, it is generally known that intensities in an X-ray powder diffraction pattern may fluctuate depending on measurement conditions. Therefore it should be understood that the 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide Form A of the present invention is not limited to the crystals that provide X-ray powder diffraction patterns identical to the X-ray powder diffraction pattern shown in Figure A, and any crystals providing X-ray powder diffraction patterns substantially the same as those shown in Figure A fall within the scope of the present invention. A person skilled in the art of X-ray powder diffraction is able to judge the substantial identity of X-ray powder diffraction patterns.

Persons skilled in the art of X-ray powder diffraction will realise that the relative intensity of peaks can be affected by, for example, grains above 30 microns in size and non-unitary aspect ratios, which may affect analysis of samples. The skilled person will also realise that the position of reflections can be affected by the precise height at which the sample sits in the diffractometer and the zero calibration of the diffractometer. The

surface planarity of the sample may also have a small effect. Hence the diffraction pattern data presented are not to be taken as absolute values. (Jenkins, R & Snyder, R.L.

'Introduction to X-Ray Powder Diffractometry' John Wiley & Sons 1996; Bunn, C.W.

(1948), Chemical Crystallography, Clarendon Press, London; Klug, H. P. & Alexander, L.

5 E. (1974), X-Ray Diffraction Procedures).

Generally, a measurement error of a diffraction angle in an X-ray powder diffractogram is about 5% or less, in particular plus or minus 0.5° 2-theta, and such degree of a measurement error should be taken into account when considering the X-ray powder diffraction pattern in Figure A and when reading Table A. Furthermore, it should be understood that intensities might fluctuate depending on experimental conditions and sample preparation (preferred orientation).

Details of Techniques Used

X-Ray Powder Diffraction

15 *Table B*

% Relative Intensity*	Definition
25 - 100	vs (very strong)
10 - 25	s (strong)
3 - 10	m (medium)
1 - 3	w (weak)

* The relative intensities are derived from diffractograms measured with fixed slits

Analytical Instrument: Siemens D5000.

The X-ray powder diffraction spectra were determined by mounting a sample of the crystalline material on a Siemens single silicon crystal (SSC) wafer mount and spreading out the sample into a thin layer with the aid of a microscope slide. The sample was spun at 30 revolutions per minute (to improve counting statistics) and irradiated with X-rays generated by a copper long-fine focus tube operated at 40kV and 40mA with a wavelength of 1.5406 angstroms. The collimated X-ray source was passed through an automatic variable divergence slit set at V20 and the reflected radiation directed through a 2mm antiscatter slit and a 0.2mm detector slit. The sample was exposed for 1 second per 0.02 degree 2-theta increment (continuous scan mode) over the range 2 degrees to 40 degrees 2-theta in theta-theta mode. The running time was 31 minutes and 41 seconds. The

- 50 -

instrument was equipped with a scintillation counter as detector. Control and data capture was by means of a Dell Optiplex 686 NT 4.0 Workstation operating with Diffract+ software. Persons skilled in the art of X-ray powder diffraction will realise that the relative intensity of peaks can be affected by, for example, grains above 30 microns in size and 5 non-unitary aspect ratios that may affect analysis of samples. The skilled person will also realise that the position of reflections can be affected by the precise height at which the sample sits in the diffractometer and the zero calibration of the diffractometer. The surface planarity of the sample may also have a small effect. Hence the diffraction pattern data presented are not to be taken as absolute values.

10

Differential Scanning Calorimetry

Analytical Instrument: Mettler DSC820e.

Typically less than 5mg of material contained in a 40 μ l aluminium pan fitted with a pierced lid was heated over the temperature range 25°C to 325°C at a constant heating rate 15 of 10°C per minute. A purge gas using nitrogen was used - flow rate 100ml per minute.

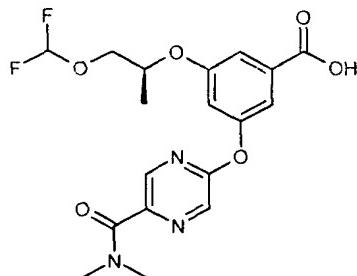
Slurrying of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide

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The X-ray powder diffraction spectra for 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide showed the initial material to be amorphous. In order to produce the crystalline form, Form A, approximately 500mg of material was placed in a vial with a magnetic follower, and approximately 2ml of Diethyl Ether added, the vial was then sealed tightly with a cap. 25 The slurry was then left to stir on a magnetic plate at ambient temperature (25°C). After 2 days, the sample was removed from the plate, the cap taken off and the slurry left to dry under ambient conditions before it was analysed by XRPD and DSC. The resulting material (Form A) was determined to be crystalline by XRPD and seen to be different from the initial amorphous material. This material (Form A) had a melting point of 75.0°C 30 (onset).

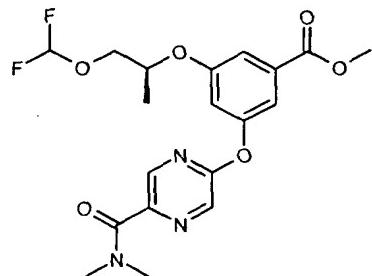
- 51 -

3-[(2S)-1-(Difluoromethoxy)propan-2-yl]oxy-5-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxy-benzoic acid



Methyl 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxy-benzoate (19.8 g, 46.6 mmol) was dissolved in THF (300 mL) and methanol (100 mL) and LiOH (1N, 51.3 mL) was added followed by water dropwise till it went cloudy. The resultant solution was stirred for 16 hours at room temperature. The organics were removed by evaporation under reduced pressure. The aqueous slurry was diluted with water (100 mL), washed with ethyl acetate (200 mL), then acidified by addition of hydrochloric acid (2N) until a solid precipitated. The resulting suspension was extracted with ethyl acetate (2 × 200 mL). The combined organic extracts were washed with water (200 mL) and brine (200 mL), dried (MgSO_4), filtered and concentrated under reduced pressure to afford the product (17.2 g, 90%). ^1H NMR δ (CDCl_3): 1.39 (d, 3H), 3.17 (s, 3H), 3.19 (s, 3H), 3.93 - 4.05 (m, 2H), 4.60 - 4.69 (m, 1H), 6.26 (t, 1H), 6.99 (t, 1H), 7.50 - 7.55 (m, 2H), 8.38 (d, 1H), 8.55 (d, 1H), 10.17 (s, 1H); m/z 412 ($\text{M}+\text{H}$)⁺

Methyl 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxy-benzoate

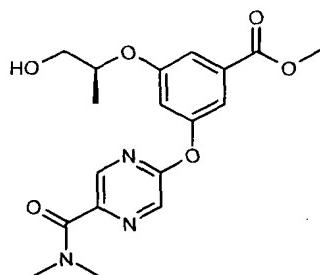


2,2-Difluoro-2-fluorosulfonyl-acetic acid (CAS no. 1717-59-5) (0.84 mL, 8.05 mmol) in acetonitrile (20 mL) was added with a syringe pump dropwise over 90 minutes to a degassed stirring mixture of methyl 3-[(dimethylcarbamoyl)pyrazin-2-yl]oxy-5-[(2S)-1-

- 52 -

hydroxypropan-2-yl]oxy-benzoate (1.5 g, 22.0 mmol) and copper (I) iodide (154 mg, 4.55 mmol) in acetonitrile (300 mL) at 55°C. The volatiles were removed under reduced pressure and the residue taken up in DCM. The mixture was filtered to and the solvent was removed under reduced pressure. The residue was purified by flash chromatography on 5 silica, eluting with a gradient of 25%-100% ethyl acetate in isohexane to afford the product (16.5 g, 62%). ¹H NMR δ (CDCl₃): 1.38 (3H, d), 3.15 (3H, s), 3.18 (3H, s), 3.91 (3H, s), 3.93 - 4.05 (2H, m), 4.62 - 4.68 (1H, m), 6.27 (1H, t), 6.95 (1H, t), 7.44 - 7.45 (1H, m), 7.49 - 7.51 (1H, m), 8.36 (1H, s), 8.53 (1H, s); *m/z* 404 (M+H⁺)

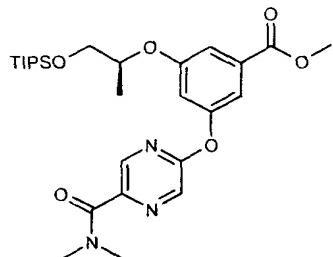
10 Methyl 3-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxy-5-[(2S)-1-hydroxypropan-2-yl]oxy-benzoate



A solution of hydrogen fluoride (70% in pyridine, 3.25 mL) was added to methyl 3-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxy-5-[(2S)-1-tripropan-2-ylsilyloxypropan-2-yl]oxy-15 benzoate in THF (300 mL) in a PTFE vessel and the resulting solution stirred for 18 hours at room temp. Further hydrogen fluoride solution (70% in pyridine, 3.25 mL) was added and the reaction stirred for an additional 66 hours. The reaction was quenched by the very careful addition of saturated aqueous sodium bicarbonate solution until the solution reached pH 8. The aqueous layer was extracted with ethyl acetate (2 × 500 mL) and the 20 combined organics dried (MgSO₄). The solvent was removed under reduced pressure and the residue was purified by flash column chromatography on silica eluting with 25 to 100% ethyl acetate in isohexane to afford the product (13.2g , 98%). ¹H NMR δ (CDCl₃) 1.32 (3H, d), 1.93 (1H, d), 3.17 (6H, d), 3.74 - 3.79 (1H, m), 3.91 (3H, s), 4.54 - 4.60 (1H, m), 6.96 (1H, t), 7.43 (1H, d), 7.51 (1H, d), 8.36 (1H, d), 8.53 (1H, d); *m/z* 376 (M+H⁺)

- 53 -

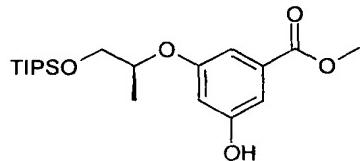
Methyl 3-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxy-5-[(2S)-1-tripropan-2-ylsilyloxypropan-2-yl]oxy-benzoate



A mixture of methyl 3-hydroxy-5-[(2S)-1-tripropan-2-ylsilyloxypropan-2-yl]oxy-benzoate (40.2 g, 105 mmol), 5-chloro-N,N-dimethyl-pyrazine-2-carboxamide (20.5 g, 110 mmol) and potassium carbonate (36.3 g, 263 mmol) in acetonitrile (500 mL) was stirred at reflux for 6 hours. The volatiles were removed under reduced pressure and ethyl acetate (500 mL) and water (500 mL) were added. The organic layer was separated, the aqueous layer was re-extracted into ethyl acetate (250 mL) and the combined organics washed with water (500 mL), brine (500 mL), dried (MgSO_4), filtered and concentrated under reduced pressure to afford the product (55.6g, 100%). ^1H NMR δ (CDCl_3) 1.01 - 1.07 (2H, m), 1.34 (3H, d), 3.14 - 3.16 (3H, s), 3.17 (3H, s), 3.72 - 3.77 (1H, m), 3.87 - 3.92 (4H, m), 4.51 (1H, m), 6.95 (1H, t), 7.39 - 7.40 (1H, m), 7.50 - 7.51 (1H, m), 8.34 (1H, d), 8.53 (1H, d); m/z 532 ($\text{M}+\text{H}^+$)

The preparation of 5-chloro-N,N-dimethyl-pyrazine-2-carboxamide was described previously.

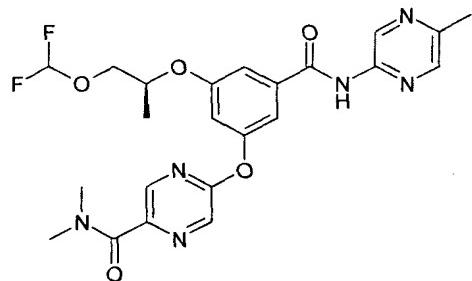
Methyl 3-hydroxy-5-[(2S)-1-tripropan-2-ylsilyloxypropan-2-yl]oxy-benzoate



To a solution of methyl 3-phenylmethoxy-5-[(2S)-1-tripropan-2-ylsilyloxypropan-2-yl]oxy-benzoate (CAS no. 871657-71-5) (47.3 g, 0.1 mol) in ethanol (500 mL) was added 10 % palladium on activated carbon (5 g) under a blanket of nitrogen. The reaction was stirred under an atmosphere of hydrogen for 16 hours. After this time the catalyst was filtered off and the solvent evaporated under reduced pressure afford the product (38.1 g,

100%). ^1H NMR δ (CDCl_3) 1.01 - 1.12 (22H, m), 1.32 (3H, d), 3.69 - 3.77 (2H, m), 3.89 (3H, s), 4.48 (1H, q), 6.62 (1H, t), 7.10 (1H, d), 7.18 (1H, t); m/z 381 (M-H $^-$)

A further alternative method for the preparation of 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide is given below:



To 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxy-benzoic acid (74.2 g, 180 mmol) was added DMF (1.4 mL, 18 mmol).

10 Dichloromethane (810 mL) and oxalyl chloride (25.2 mL, 289 mmol) were charged and the reaction left to stir at ambient temperature for 2 hours. The solvent was evaporated under reduced pressure, azeotroped with toluene (2 × 600 mL) and the resulting oil dissolved in pyridine (392ml) and dichloromethane (500ml).

15 A solution of 5-methylpyrazin-2-amine (CAS no. 5521-58-4) (29.7 g, 272 mmol) in pyridine (549 mL) was charged dropwise to the stirred solution and the reaction stirred at ambient temperature for 20 hours. The solvent was evaporated under reduced pressure and the residue was taken up into ethyl acetate (1200 mL), washed with water (1200 mL), 1M citric acid (2 × 780 mL), saturated aqueous sodium hydrogen carbonate (2 × 780 mL), saturated aqueous brine (780 mL) and the combined organic extracts were dried (MgSO_4) and evaporated under reduced pressure the residue was purified by flash chromatography to afford the title compound (66 g).

20 To a sample of this material (64 g, 127 mmol) was added diethyl ether (640 mL) and the resulting slurry stirred overnight.

A solid was filtered, washed with diethyl ether (320 mL) and dried under vacuum at ambient temperature overnight to provide a white crystalline solid (56g).

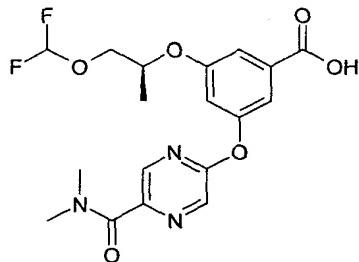
25 This material had an X-ray powder diffraction pattern consistent with that described for 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-

- 55 -

yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide Form A described previously. ¹H NMR and mass spectrometry data were consistent with those described previously.

- 5 The preparation of 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxy-benzoic acid is described below.

3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxy-benzoic acid

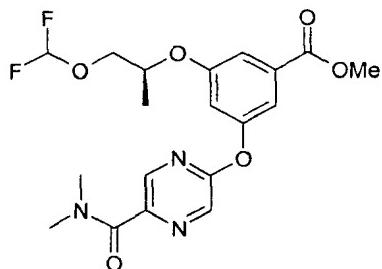


10 To methyl 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxy-benzoate (76.3 g, 179 mmol) was added NMP (534 mL), water (305 mL) and the solution was stirred at 0 °C. 2N sodium hydroxide (152 mL, 305 mmol) was charged dropwise and the reaction stirred for 4 hours. Acetic acid (41 mL, 718 mmol) was charged 15 dropwise followed by water (1068 mL) and 1N HCl (400 mL) was added until pH 3 was reached and some material oiled out. The aqueous layer was extracted with toluene (3 × 988 mL) and combined with the material that oiled out which was dissolved in ethyl acetate (988 mL) and the combined organic layers were washed with water (988 mL), saturated aqueous brine (988 mL), dried (MgSO_4) and evaporated under reduced pressure. 20 The residue was purified by flash chromatography eluting with 5% MeOH in dichloromethane to afford the title compound (64 g).

¹H NMR δ (DMSO-d₆) 1.27 (3H, d), 3.03 (6H, s), 3.99 - 4.04 (2H, m), 4.76 - 4.80 (1H, m), 6.52-6.91 (1H, t), 7.21 (1H, t), 7.34 - 7.38 (2H, m), 8.42 (1H, d), 8.56 (1H, d)

- 56 -

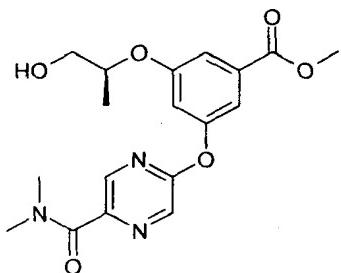
Methyl 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxy-benzoate



5 A solution of methyl 3-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxy-5-[(2S)-1-hydroxypropan-2-yl]oxy-benzoate (94 g, 250 mmol) in acetonitrile (1130 mL) was degassed with nitrogen then copper(I) iodide (9.54 g, 50 mmol) was charged and the solution was heated to 55°C. A solution of 2,2-difluoro-2-fluorosulfonyl-acetic acid (CAS no. 1717-59-5) (46.6 mL, 450 mmol) in acetonitrile (188 mL) was charged dropwise. After 3 hours the solvent was evaporated under reduced pressure at 25 °C. The residue was taken up in dichloromethane (500 mL) and filtered. The solid was washed further with dichloromethane until the washings were clear. The solvent was evaporated under reduced pressure at 25 °C and the residue was purified by flash chromatography eluting with 100% ethyl acetate to afford the title compound (54 g).

10 15 ¹H NMR δ (DMSO-d₆) 1.28 (3H, d), 2.99 - 3.08 (6H, m), 3.86 (3H, s), 3.98 - 4.07 (2H, m), 4.78 - 4.82 (1H, m), 6.50-6.90 (1H, t), 7.25 (1H, t), 7.38 - 7.40 (2H, m), 8.42 (1H, d), 8.55 (1H, d)

20 Methyl 3-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxy-5-[(2S)-1-hydroxypropan-2-yl]oxy-benzoate



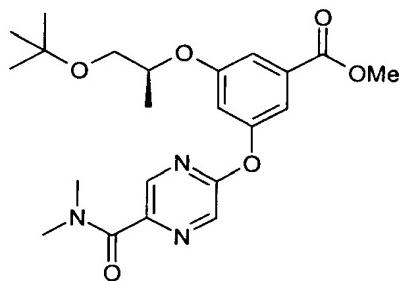
A solution of methyl 3-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxy-5-[(2S)-1-[(2-methylpropan-2-yl)oxy]propan-2-yl]oxy-benzoate (170 g, 0.39 mol) in formic acid

(850 mL) was heated to 90 °C for 3 hours. Ethyl acetate (1700 mL), water (1700 mL) and saturated aqueous brine (850 mL) was added and the aqueous layer separated and extracted with ethyl acetate (850 mL) and the combined organic layers were washed with saturated aqueous brine (850 mL), dried (MgSO_4) and evaporated under reduced pressure. The residue was dissolved with ethyl acetate (1500 mL), water (1500 mL) and methanol (150 mL). Sodium carbonate (170 g) was added and the biphasic solution heated to reflux for 2 hours. The aqueous layer was separated and the organic layer washed with water (1700 mL). The combined aqueous phases were extracted with ethyl acetate (850 mL) and the combined organic layers dried (MgSO_4) and evaporated under reduced pressure. The residue was purified by flash chromatography eluting with 100% ethyl acetate to obtain a the title compound (148 g).

¹H NMR δ (DMSO-d₆) 1.23 (3H, d), 3.04 (6H, s), 3.47 - 3.56 (2H, m), 3.86 (3H, s), 4.49 - 4.53 (1H, m), 4.86 (1H, t), 7.19 (1H, t), 7.34 - 7.35 (1H, m), 7.38 - 7.39 (1H, m), 8.42 (1H, d), 8.55 (1H, d)

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Methyl 3-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxy-5-[(2S)-1-[(2-methylpropan-2-yl)oxy]propan-2-yl]oxy-benzoate



To methyl 3-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxy-5-hydroxy-benzoate (10 g,

20 32 mmol) was added triphenylphosphine (10.3 g, 39.4 mmol), THF (100 mL) and (2R)-1-[(2-methylpropan-2-yl)oxy]propan-2-ol (CAS no. 136656-73-0) (5.21 g, 39.4 mmol). The resulting slurry was cooled to 0 °C and diethylazodicarboxylate (50% w/v in toluene, 13.7 mL, 39.4 mmol) was charged dropwise keeping the temperature below 10 °C. After 2 hours the solvent was evaporated under reduced pressure and taken up with ethyl acetate (23 mL). A solid crashed out which was filtered and the mother liquors were evaporated under reduced pressure, taken up into ethyl acetate (23 mL) and isohexane (53 mL) and the resulting solid filtered, the mother liquors were evaporated under reduced pressure and the

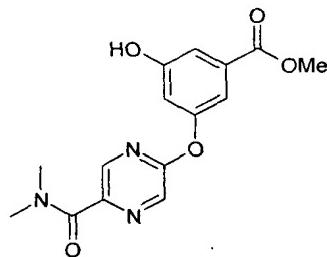
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- 58 -

resulting residue was purified by flash chromatography eluting with 80% ethyl acetate / 20% isohexane to afford the product (13.5 g).

¹H NMR δ (DMSO-d₆) 1.12 (9H, s), 1.25 (3H, d), 3.03 (6H, s), 3.41 - 3.45 (1H, m), 3.47 - 3.51 (1H, m), 3.85 (3H, s), 4.55 - 4.57 (1H, m), 7.20 (1H, t), 7.34 - 7.35 (1H, m), 7.40 - 7.41 (1H, m), 8.41 (1H, d), 8.54 (1H, d)

Methyl 3-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxy-5-hydroxy-benzoate



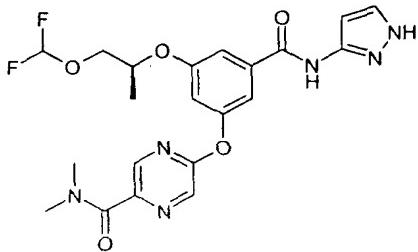
To methyl 3,5-dihydroxybenzoate (CAS no. 2150-44-9) (85 g, 0.49 mol) was added 5-chloro-N,N-dimethyl-pyrazine-2-carboxamide (88.9 g, 0.48 mol), DMSO (1000 mL,) and cesium carbonate (418 g, 1.2 mol) and the mixture was heated to 50 °C for 3 hours. Water (1577 mL,) was charged followed by diethyl ether (540 mL). To the aqueous layer was charged 5M hydrochloric acid solution (395 mL, 1.97 mol) and the resulting white solid was filtered washed with water (2 × 311 mL) and dried under vacuum at 40 °C over P₂O₅ overnight to give the desired compound (143 g).

¹H NMR δ (DMSO-d₆) 3.03 (6H, s), 3.84 (3H, s), 6.92 (1H, t), 7.21 - 7.22 (1H, m), 7.28 - 7.29 (1H, m), 8.41 (1H, d), 8.53 (1H, d), 10.20 (1H, s).

The preparation of 5-chloro-N,N-dimethyl-pyrazine-2-carboxamide was described previously.

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Example 3: 5-[3-[(2S)-1-(Difluoromethoxy)propan-2-yl]oxy-5-(1H-pyrazol-3-ylcarbamoyl)phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide



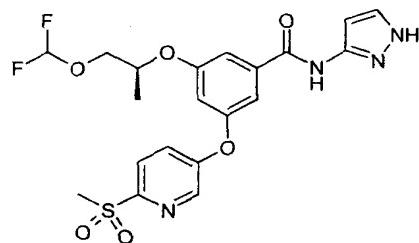
- 59 -

Trifluoroacetic acid (2 mL) was added to a solution of tert-butyl 3-[[2S]-1-(difluoromethoxy)propan-2-yl]oxy-5-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxybenzoyl]amino]pyrazole-1-carboxylate (150 mg, 0.26 mmol) in DCM (16 mL) and stirred at RT for 2 hours. The solvent was removed *in vacuo*, DCM (20 mL) added and the mixture washed with water (20 mL), saturated sodium bicarbonate solution (20 mL), brine (20 mL), dried (MgSO_4) and reduced *in vacuo* to give the desired compound (94 mg).

^1H NMR δ (CDCl_3): 1.38 (d, 3H), 3.15 (s, 3H), 3.18 (s, 3H), 3.95 - 4.03 (m, 2H), 4.62 - 4.69 (m, 1H), 6.25 (t, 1H), 6.84 (s, 1H), 6.92 (t, 1H), 7.31 (s, 1H), 7.37 (s, 1H), 7.44 (d, 1H), 8.39 (d, 1H), 8.49 (d, 1H), 9.71 (s, 1H), 10.04 (s, 1H); m/z 477 ($\text{M}+\text{H}$)⁺

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Example 4 - 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-(6-methylsulfonylpyridin-3-yl)oxy-N-(1H-pyrazol-3-yl)benzamide

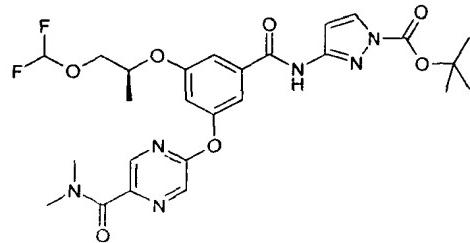


The following compound was prepared in an analogous fashion to example 3 from tert-butyl 3-[[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-(6-methylsulfonylpyridin-3-yl)oxy-benzoyl]amino]pyrazole-1-carboxylate.

^1H NMR δ (CDCl_3): 1.39 (d, 3H), 3.23 (s, 3H), 3.95 - 4.04 (m, 2H), 4.64 - 4.71 (m, 1H), 6.26 (t, 1H), 6.83 - 6.86 (m, 2H), 7.19 (s, 1H), 7.35 (s, 1H), 7.45 - 7.48 (m, 1H), 7.52 (s, 1H), 8.07 (d, 1H), 8.48 (d, 1H), 8.65 (s, 1H); m/z 483 ($\text{M}+\text{H}$)⁺

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tert-Butyl 3-[[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-(6-methylsulfonylpyridin-3-yl)oxy-benzoyl]amino]pyrazole-1-carboxylate



- 60 -

1-Chloro-N,N,2-trimethyl-prop-1-en-1-amine (0.11 mL, 0.80 mmol) was added to a solution of 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxy-benzoic acid (0.22 g, 0.53 mmol) in DCM (5 mL) and stirred for 1 hour. tert-Butyl 3-aminopyrazole-1-carboxylate (CAS no. 863504-94-1) (147 mg, 0.80 mmol) then pyridine (0.09 mL, 1.07 mmol) were added and the reaction stirred for a further 45 minutes before being reduced *in vacuo* and partitioned between ethyl acetate (50 mL) and water (50 mL). The aqueous layer was further extracted into ethyl acetate (50 mL) and the combined organics washed with water (50 mL), brine (50 mL), dried (MgSO_4), and reduced *in vacuo*. The crude residue was chromatographed on silica, eluting with 20-50% ethyl acetate in isohexane, to give the desired compound (0.15 g).

¹H NMR δ (CDCl_3): 1.38 (d, 3H), 1.60 (s, 9H), 3.16 (s, 3H), 3.19 (s, 3H), 3.93 - 4.04 (m, 2H), 4.60 - 4.64 (m, 1H), 6.26 (t, 1H), 6.95 (t, 1H), 7.09 (d, 1H), 7.27 - 7.28 (m, 1H), 7.34 (t, 1H), 8.01 (d, 1H), 8.37 (d, 1H), 8.53 (d, 1H), 8.97 (s, 1H); *m/z* 577 ($\text{M}+\text{H}$)⁺

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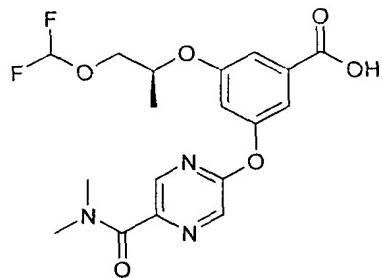
tert-Butyl 3-[[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-(6-methylsulfonylpyridin-3-yl)oxy-benzoyl]amino]pyrazole-1-carboxylate, used in the preparation of Example 4, was prepared in an analogous fashion from 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-(6-methylsulfonylpyridin-3-yl)oxy-benzoic acid.

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Structure	<i>m/z</i>	NMR
	583 ($\text{M}+\text{H}$) ⁺	¹ H NMR δ (CDCl_3): 1.39 (d, 3H), 1.64 (s, 9H), 3.24 (s, 3H), 3.95 - 4.04 (m, 2H), 4.64 - 4.68 (m, 1H), 6.26 (t, 1H), 6.85 (t, 1H), 7.07 (d, 1H), 7.14 (t, 1H), 7.30 (t, 1H), 7.46 - 7.48 (m, 1H), 8.01 (d, 1H), 8.09 (d, 1H), 8.48 (d, 1H), 8.67 (s, 1H)

- 61 -

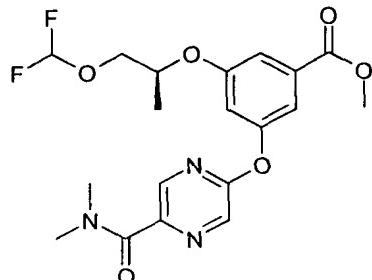
3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxybenzoic acid



Lithium hydroxide monohydrate (45 mg, 1.06 mol) in water (5 mL) was added to a solution of methyl 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxybenzoate (0.3 g, 0.71 mmol) in THF (10 mL) and stirred at RT for 20 hours. The THF was removed *in vacuo* and the aqueous layer was washed with ethyl acetate (50 mL) to remove any impurities. The aqueous layer was acidified and extracted into ethyl acetate (2 × 50 mL) then the combined organics washed with brine (50 mL), dried (MgSO_4) and the solvent removed *in vacuo* to give the desired compound (0.22 g).

^1H NMR δ (CDCl_3): 1.39 (d, 3H), 3.17 (s, 3H), 3.19 (s, 3H), 3.93 - 4.05 (m, 2H), 4.60 - 4.69 (m, 1H), 6.26 (t, 1H), 6.99 (t, 1H), 7.50 - 7.55 (m, 2H), 8.38 (d, 1H), 8.55 (d, 1H), 10.17 (s, 1H); m/z 412 ($\text{M}+\text{H}^+$)⁺

Methyl 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[5-(dimethylcarbamoyl)pyrazin-2-yl]oxybenzoate



A mixture of methyl 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-hydroxybenzoate (0.25 g, 0.91 mmol), 5-chloro-N,N-dimethyl-pyrazine-2-carboxamide (168 mg, 0.91 mmol) and potassium carbonate (250 mg, 1.81 mmol) in acetonitrile (5 mL) was stirred in a microwave reactor at 140°C for 5 hours. The mixture was reduced *in vacuo* and ethyl acetate (50 mL) added. The mixture was washed with water (50 mL), brine (50 mL), dried

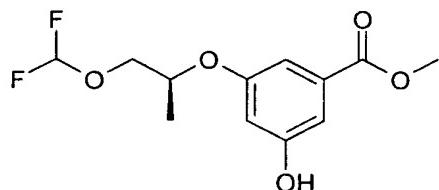
- 62 -

(MgSO₄) and reduced *in vacuo*. The residue was chromatographed on silica, eluting with 20 to 70% ethyl acetate in isohexane, to give the desired compound (0.3 g).

¹H NMR δ (CDCl₃): 1.38 (d, 3H), 3.15 (s, 3H), 3.18 (s, 3H), 3.91 (s, 3H), 3.93 - 4.04 (m, 2H), 4.61 - 4.69 (m, 1H), 6.26 (t, 1H), 6.96 (t, 1H), 7.44 - 7.45 (m, 1H), 7.50 - 7.51 (m, 1H), 8.36 (d, 1H), 8.53 (d, 1H); *m/z* 426 (M+H)⁺

The preparation of 5-chloro-N,N-dimethyl-pyrazine-2-carboxamide was described earlier.

Methyl 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-hydroxy-benzoate



10

Methyl 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-phenylmethoxy-benzoate (0.48 g, 1.1 mmol) was dissolved in ethanol (10 mL) and THF (10 mL) and the flask evacuated and purged with argon (3 times). 10% Palladium on carbon (140 mg) was added and the flask further evacuated and finally purged with hydrogen gas. The reaction mixture was stirred at RT for 20 hours until completion. The reaction mixture was evacuated and purged with argon (3 times) then the catalyst removed by filtration through celite®. The filtrate was concentrated *in vacuo* to give the desired compound (1.05 g).

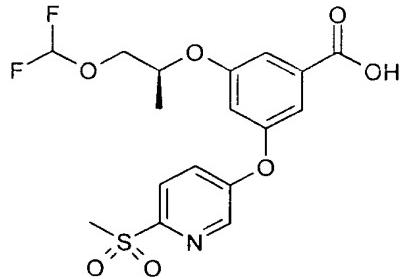
¹H NMR δ (CDCl₃): 1.35 (d, 3H), 3.90 (s, 3H), 3.90 - 4.02 (m, 2H), 4.57 - 4.64 (m, 1H), 5.20 (s, 1H), 6.26 (t, 1H), 6.63 (t, 1H), 7.14 - 7.15 (m, 1H), 7.17 - 7.18 (m, 1H); *m/z* 275 (M-H)⁻

The preparation of methyl 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-phenylmethoxy-benzoate was described earlier.

25 The preparation of 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-(6-methylsulfonylpyridin-3-yl)oxy-benzoic acid is described below.

- 63 -

3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-(6-methylsulfonylpyridin-3-yl)oxy-benzoic acid



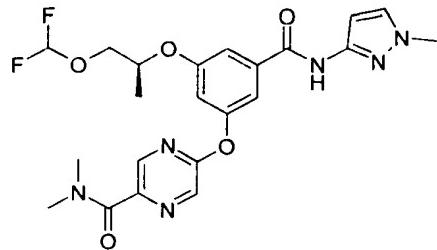
A mixture of methyl 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-hydroxy-benzoate (233 mg, 0.84 mmol), 5-bromo-2-methylsulfonyl-pyridine (CAS no. 98626-95-0) (200 mg, 0.84 mmol), cesium carbonate (549 mg, 1.69 mmol) and bromotris(triphenylphosphine)copper(I) (157 mg, 0.17 mmol) in DMA (5 mL) was stirred in a microwave reactor at 160°C for 6 hours. Ethyl acetate (50 mL) and water were added and the aqueous layer was acidified and extracted with ethyl acetate (2 × 50 mL). The combined organics were washed with brine, dried (MgSO_4) and reduced *in vacuo* to give the desired compound (0.16 g).

^1H NMR δ (d_6 -DMSO): 1.28 (d, 3H), 3.27 (s, 3H), 3.95 - 4.04 (m, 2H), 4.78 - 4.85 (m, 1H), 6.71 (t, 1H), 7.14 - 7.16 (m, 1H), 7.22 - 7.23 (m, 1H), 7.37 - 7.40 (m, 1H), 7.66 - 7.70 (m, 1H), 8.06 (d, 1H), 8.61 (d, 1H), 12.85 (s, 1H); m/z 418 ($\text{M}+\text{H}$)⁺

15

The preparation of methyl 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-hydroxy-benzoate was described earlier.

Example 5: 5-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(1-methylpyrazol-3-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide



A mixture of 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-hydroxy-N-(1-methylpyrazol-3-yl)benzamide (0.1 g, 0.29 mmol), 5-chloro-N,N-dimethyl-pyrazine-2-carboxamide

- 64 -

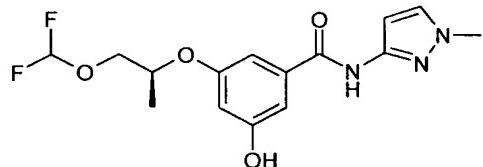
(66 mg, 0.35 mmol) and potassium carbonate (81 mg, 0.59 mmol) in acetonitrile (5 mL) was stirred in a microwave reactor at 160°C for 6 hours. The resulting mixture was reduced *in vacuo* and ethyl acetate (50 mL) added. The organics were washed with water (50 mL), brine (50 mL), dried (MgSO_4), filtered and reduced *in vacuo*. The residue was chromatographed on silica, eluting with 10-100% ethyl acetate in isohexane to give the desired compound (52 mg).

¹H NMR δ (CDCl_3): 1.30 (d, 3H), 3.08 (s, 3H), 3.11 (s, 3H), 3.69 (s, 3H), 3.85 - 3.97 (m, 2H), 4.56 (sextet, 1H), 6.18 (t, 1H), 6.73 (d, 1H), 6.85 (t, 1H), 7.19 - 7.21 (m, 2H), 7.27 - 7.29 (m, 1H), 8.29 (d, 1H), 8.45 (d, 1H), 8.76 (s, 1H); m/z 491 ($\text{M}+\text{H}$)⁺

10

The synthesis of 5-chloro-N,N-dimethyl-pyrazine-2-carboxamide was described previously.

3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-hydroxy-N-(1-methylpyrazol-3-yl)benzamide



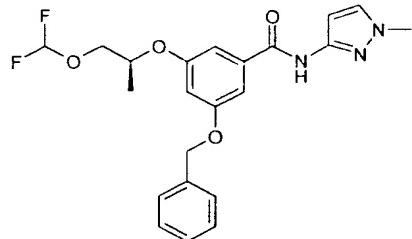
3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-N-(1-methylpyrazol-3-yl)-5-phenylmethoxybenzamide (0.1 g, 0.23 mmol) was dissolved in ethanol (3 mL) and THF (3 mL) and the flask evacuated and purged with argon (3 times). 10% Palladium on carbon (0.01 g) was added and the flask further evacuated and finally purged with hydrogen gas. The reaction mixture was stirred at RT for 20 hours until completion. The reaction mixture was evacuated and purged with nitrogen (3 times). The catalyst was filtered off through celite and the filtrate concentrated *in vacuo* to give the desired compound (70 mg).

¹H NMR δ (CDCl_3): 1.28 (d, 3H), 3.71 (s, 3H), 3.80-3.95 (m, 2H), 4.51 (sextet, 1H), 5.96-6.36 (t, 1H), 6.53 (s, 1H), 6.73 (s, 1H), 6.91 (s, 1H), 6.96 (s, 1H), 7.22 (s, 1H), 8.83 (s, 1H); m/z 342 ($\text{M}+\text{H}$)⁺

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- 65 -

3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-N-(1-methylpyrazol-3-yl)-5-phenylmethoxy-benzamide

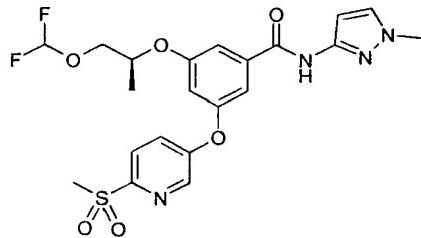


DIPEA (0.198 mL, 1.14 mmol) was added to a mixture of 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-phenylmethoxy-benzoic acid (0.10 g, 0.28 mmol), 1-methylpyrazol-3-amine (CAS no. 1904-31-0) (39 mg, 0.4 mmol) and HATU (0.227 g, 0.6 mmol) in DMF (3 mL) and stirred at RT for 20 hours. Ethyl acetate (30 mL) was added and the mixture washed with water (30 mL), brine (30 mL), dried (MgSO_4), filtered and reduced *in vacuo*. The residue was chromatographed on silica, eluting with a gradient of 0-100% ethyl acetate in isohexane, to give the desired compound (0.1 g).

^1H NMR δ (CDCl_3): 1.36 (d, 3H), 3.68 (s, 3H), 3.82-3.95 (m, 2H), 4.48 (sextet, 1H), 5.00 (s, 2H), 6.19 (t, 1H), 6.63 (s, 1H), 6.73 (s, 1H), 6.93 (s, 1H), 7.03 (s, 1H), 7.28 (m, 1H), 7.35 (m, 5H), 8.59 (s, 1H); m/z 432 ($\text{M}+\text{H})^+$

15 The synthesis of 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-phenylmethoxy-benzoic acid was described earlier.

Example 6: 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-N-(1-methylpyrazol-3-yl)-5-(6-methylsulfonylpyridin-3-yl)oxy-benzamide



20 A mixture of 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-hydroxy-N-(1-methylpyrazol-3-yl)benzamide (100 mg, 0.29 mmol), 5-bromo-2-methylsulfonyl-pyridine (CAS no. 98626-95-0) (77 mg, 0.32 mmol), cesium carbonate (191 mg, 0.59 mmol) and bromotris(triphenylphosphine)copper(I) (55 mg, 0.06 mmol) in DMA (5 mL) was stirred

- 66 -

in a microwave reactor at 160°C for 6 hours. Ethyl acetate (50 mL) was added and washed with water (50 mL), brine (50 mL), dried (MgSO_4), filtered and reduced *in vacuo*. The residue was chromatographed on silica, eluting with 10-80% ethyl acetate in isohexane to give the desired compound (31 mg).

5 ^1H NMR δ (CDCl_3): 1.30 (d, 3H), 3.16 (s, 3H), 3.72 (s, 3H), 3.85 - 3.95 (m, 2H), 4.53 - 4.59 (m, 1H), 6.18 (t, 1H), 6.71 (d, 1H), 6.75 (t, 1H), 7.09 (t, 1H), 7.22 (d, 1H), 7.25 (t, 1H), 7.37 - 7.39 (m, 1H), 7.99 (d, 1H), 8.39 (d, 1H), 8.62 (s, 1H); m/z 495 (M-H) $^-$

10 The synthesis of 3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-hydroxy-N-(1-methylpyrazol-3-yl)benzamide was described previously.

BIOLOGICAL

Tests:

The biological effects of the compounds of formula (I) may be tested in the following way:

15

(1) Enzymatic activity

Enzymatic activity of recombinant human pancreatic GLK may be measured by incubating GLK, ATP and glucose. The rate of product formation may be determined by coupling the assay to a G-6-P dehydrogenase, NADP/NADPH system and measuring the linear increase with time of optical density at 340nm Brocklehurst et al (Diabetes 2004, 53, 535-541). Activation of GLK by compounds can be assessed using this assay in the presence or absence of GLKRP as described in Brocklehurst et al (Diabetes 2004, 53, 535-541).

Compounds of the invention were assessed in the absence of GLKRP as described by

25

Brocklehurst et al and activated glucokinase with EC₅₀ values as shown below.

Table C

Example Number	EC ₅₀ Value (μM)
1	0.069
2	0.055
3	0.065
4	0.033

5	0.079
6	0.077

Production of recombinant GLK and GLKRP:

Human GLK and GLKRP cDNA was obtained by PCR from human pancreatic and hepatic mRNA respectively, using established techniques described in Sambrook J, Fritsch EF & Maniatis T, 1989. PCR primers were designed according to the GLK and GLKRP cDNA sequences shown in Tanizawa *et al.*, Proc Natl Acad Sci 1991 Aug 15;88(16):7294-7 and Warner *et al.*, Mamm Genome, 1995 Aug;6(8):532-6.

Cloning in Bluescript II vectors

GLK and GLKRP cDNA was cloned in E. coli using pBluescript II,

Transformations

E. Coli transformations were generally carried out by electroporation. 400 mL cultures of strains DH5a or BL21(DE3) were grown in L-broth to an OD 600 of 0.5 and harvested by centrifugation at 2,000g. The cells were washed twice in ice-cold deionised water, resuspended in 1mL 10% glycerol and stored in aliquots at -70°C. Ligation mixes were desalted using Millipore V series™ membranes (0.0025mm pore size). 40mL of cells were incubated with 1mL of ligation mix or plasmid DNA on ice for 10 minutes in 0.2cm electroporation cuvettes, and then pulsed using a Gene Pulser™ apparatus (BioRad) at 0.5kVcm⁻¹, 250mF. Transformants were selected on L-agar supplemented with tetracycline at 10mg/mL or ampicillin at 100mg/mL.

Expression

GLK was expressed from the vector pTB375NBSE in E.coli BL21 cells,, producing a recombinant protein containing a 6-His tag immediately adjacent to the N-terminal methionine. Alternatively, another suitable vector is pET21(+)DNA, Novagen, Cat number 697703. The 6-His tag was used to allow purification of the recombinant protein on a column packed with nickel-nitrilotriacetic acid agarose purchased from Qiagen (cat no 30250).

GLKRP was expressed from the vector pFLAG CTC (IBI Kodak) in E.coli BL21 cells, producing a recombinant protein containing a C-terminal FLAG tag. The protein was purified initially by DEAE Sepharose ion exchange followed by utilisation of the FLAG tag for final purification on an M2 anti-FLAG immunoaffinity column purchased from Sigma-Aldrich (cat no. A1205).

(2) Oral Glucose Tolerance Test (OGTT)

Oral glucose tolerance tests (G.J Coope *et al*, British Journal of Pharmacology, (2006) 149, 328-335) may be performed on conscious Zucker obese fa/fa rats (age 12-13 weeks or older) fed a high fat diet (45 % kcal fat) for at least two weeks prior to experimentation. The animals are fasted for 2 hours before use for experiments. A test compound or a vehicle is given orally 120 minutes before oral administration of a glucose solution at a dose of 2 g/kg body weight. Blood glucose levels are measured using a Accuchek glucometer from tail bled samples taken at different time points before and after administration of glucose (time course of 60 minutes). A time curve of the blood glucose levels is generated and the area-under-the-curve (AUC) for 120 minutes calculated (the time of glucose administration being time zero). Percent reduction in glucose excursion is determined using the AUC in the vehicle-control group as zero percent reduction.

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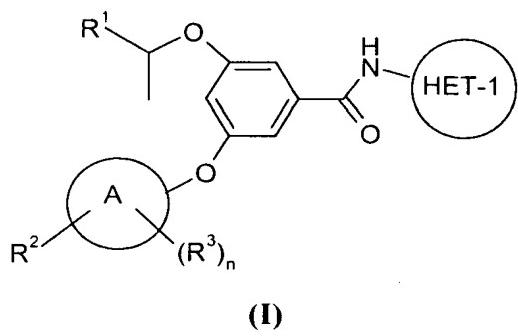
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Claims:

1. A compound of Formula (I):



wherein:

R¹ is selected from fluoromethoxymethyl, difluoromethoxymethyl and trifluoromethoxymethyl;

10 R² is a substituent on a carbon atom of Ring A and is selected from -C(O)NR⁴R⁵, -SO₂NR⁴R⁵, -S(O)_pR⁴ and HET-2;

HET-1 is a 5- or 6-membered, C-linked heteroaryl ring containing a nitrogen atom in the 2-position relative to the amide nitrogen to which the ring is attached and optionally 1 or 2 further ring heteroatoms independently selected from O, N and S; which ring is optionally substituted on an available carbon atom, or on a ring nitrogen atom provided it is not thereby quaternised, with 1 or 2 substituents independently selected from R⁶;

15 HET-2 is a 4-, 5- or 6-membered, C- or N-linked heterocyclyl ring containing 1, 2, 3 or 4 heteroatoms independently selected from O, N and S, wherein a -CH₂- group can optionally be replaced by a -C(O)-, and wherein a sulphur atom in the heterocyclic ring
20 may optionally be oxidised to a S(O) or S(O)₂ group, which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁷;

R³ is a substituent on a carbon atom of Ring A and is selected from halo;

25 R⁴ is selected from hydrogen, (1-4C)alkyl [optionally substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and -C(O)NR⁵R⁵], (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and HET-2;

R⁵ is hydrogen or (1-4C)alkyl;

R⁶ is independently selected from (1-4C)alkyl, hydroxy(1-4C)alkyl, (1-4C)alkoxy(1-4C)alkyl, (1-4C)alkylS(O)p(1-4C)alkyl, amino(1-4C)alkyl, (1-4C)alkylamino(1-4C)alkyl, di(1-4C)alkylamino(1-4C)alkyl, and/or (for R⁶ as a substituent on carbon) halo;

5 R⁷ is selected from (1-4C)alkyl, -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, (1-4C)alkoxy(1-4C)alkyl, hydroxy(1-4C)alkyl, -S(O)pR⁵ and/or (for R⁷ as a substituent on carbon) hydroxy and (1-4C)alkoxy;

Ring A is a 5- or 6- membered heteroaryl ring, containing 1, 2 or 3 ring heteroatoms independently selected from O, S and N; which ring optionally further substituted on an

10 available nitrogen atom (provided it is not thereby quaternised) by a substituent selected from R⁸;

R⁸ is selected from (1-4C)alkyl, (3-6C)cycloalkyl, hydroxy(1-4C)alkyl, (1-4C)alkoxy(1-4C)alkyl, -C(O)(1-4C)alkyl, benzyl, and (1-4C)alkylsulfonyl;

p is (independently at each occurrence) 0, 1 or 2;

15 n is 0, 1 or 2;

or a salt thereof.

2. A compound of the formula (I) as claimed in Claim 1 or a salt thereof wherein R¹ is fluoromethoxymethyl or difluoromethoxymethyl.

20

3. A compound of the formula (I) as claimed in Claim 1 or Claim 2 or a salt thereof wherein R¹ has the (S) configuration.

4. A compound of the formula (I) as claimed in Claim 1, which is any one of the

25 following:

3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-N-(5-methylpyrazin-2-yl)-5-(6-methylsulfonylpyridin-3-yl)oxy-benzamide;

5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide;

- 73 -

5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-(1H-pyrazol-3-ylcarbamoyl)phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide;
3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-(6-methylsulfonylpyridin-3-yl)oxy-N-(1H-pyrazol-3-yl)benzamide;
5 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(1-methylpyrazol-3-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide;
3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-N-(1-methylpyrazol-3-yl)-5-(6-methylsulfonylpyridin-3-yl)oxy-benzamide;
or a salt thereof.

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5. A pharmaceutical composition comprising a compound according to any one of Claims 1 to 4, or a pharmaceutically-acceptable salt thereof, together with a pharmaceutically acceptable diluent or carrier.

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6. A compound according to any one of Claims 1 to 4 or a pharmaceutically-acceptable salt thereof for use as a medicament.

7. The use of a compound according to any one of Claims 1 to 4, or a pharmaceutically-acceptable salt thereof for the preparation of a medicament for treatment 20 of a disease mediated through GLK.

8. The use of a compound according to any one of Claims 1 to 4, or a pharmaceutically-acceptable salt thereof for the preparation of a medicament for treatment of type 2 diabetes.

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9. A method of treating GLK mediated diseases by administering an effective amount of a compound of Formula (I) as claimed in any one of Claims 1 to 4 or a pharmaceutically-acceptable salt thereof, to a mammal in need of such treatment.

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10. The method of Claim 9 wherein the GLK mediated disease is type 2 diabetes.

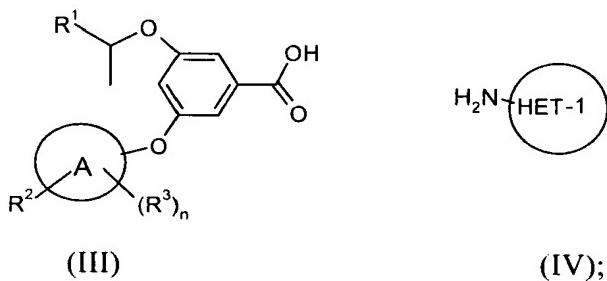
- 74 -

11. A compound according to any one of Claims 1 to 4 or a pharmaceutically-acceptable salt thereof for use as a medicament for the treatment of a disease mediated through GLK.

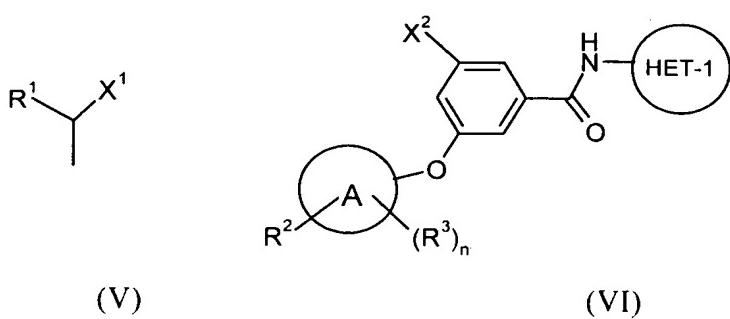
5 12. A compound according to Claim 11 wherein the disease mediated through GLK is type-2 diabetes.

13. A process for the preparation of a compound of Formula (I) as claimed in any one of Claims 1 to 4, which comprises a process a) to e) (wherein the variables are as defined for compounds of Formula (I) in Claim 1 unless otherwise stated):

(a) reaction of an acid of Formula (III) or activated derivative thereof with a compound of Formula (IV), wherein R¹ is as defined for formula (I) or is a precursor thereof;

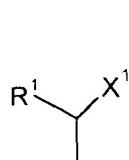


(b) reaction of a compound of Formula (V) with a compound of Formula (VI)

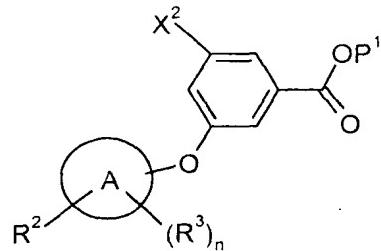


wherein X^1 is a leaving group and X^2 is a hydroxyl group or X^1 is a hydroxyl group and X^2 is a leaving group, and wherein R^1 is as defined for formula (I) or is a precursor thereof; process (b) could also be accomplished using the intermediate ester Formula (VII), wherein P^1 is a protecting group as hereinafter described, followed by ester hydrolysis and amide formation by procedures described elsewhere and well known to those skilled in the art;

- 75 -



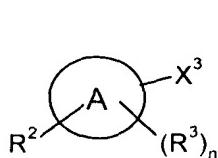
(V)



(VII)

or

(c) reaction of a compound of Formula (VIII) with a compound of Formula (IX)



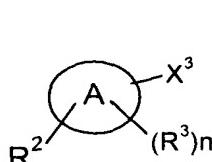
(VIII)



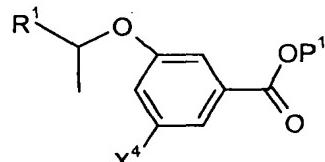
(IX)

wherein X^3 is a leaving group or an organometallic reagent and X^4 is a hydroxyl group or X^3 is a hydroxyl group and X^4 is a leaving group or an organometallic reagent, and wherein R^1 is as defined for formula (I) or is a precursor thereof;

process (c) could also be accomplished using the intermediate ester Formula (X), followed by ester hydrolysis and amide formation by procedures described elsewhere and well known to those skilled in the art;



(VIII)

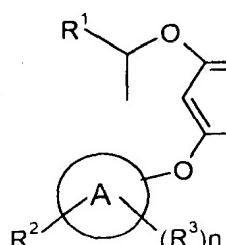


(X)

15

or

(d) reaction of a compound of Formula (XI) with a compound of Formula (XII),



(XI)

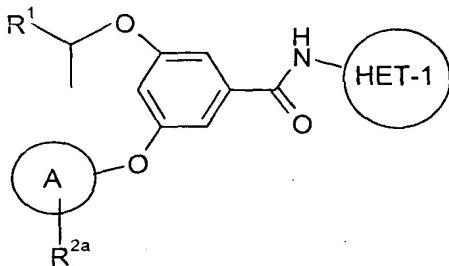


(XII);

- 76 -

wherein X^5 is a leaving group; and wherein R^1 is as defined for formula (I) or is a precursor thereof; or

e) reaction of a compound of formula (XIII)



(XIII)

wherein R^{2a} is a precursor to R^2 as $-CONR^4R^5$ or $-SO_2R^4R^5$, such as a carboxylic acid, ester or anhydride (for $R^2 = -CONR^4R^5$) or the sulfonic acid equivalents (for R^2 is $-SO^2NR^4R^5$); with an amine of formula $-NR^4R^5$;

and thereafter, if necessary:

- i) converting a compound of Formula (I) into another compound of Formula (I);
- ii) converting a precursor of R^1 into R^1 ;
- iii) removing any protecting groups; and/or
- iv) forming a salt thereof.

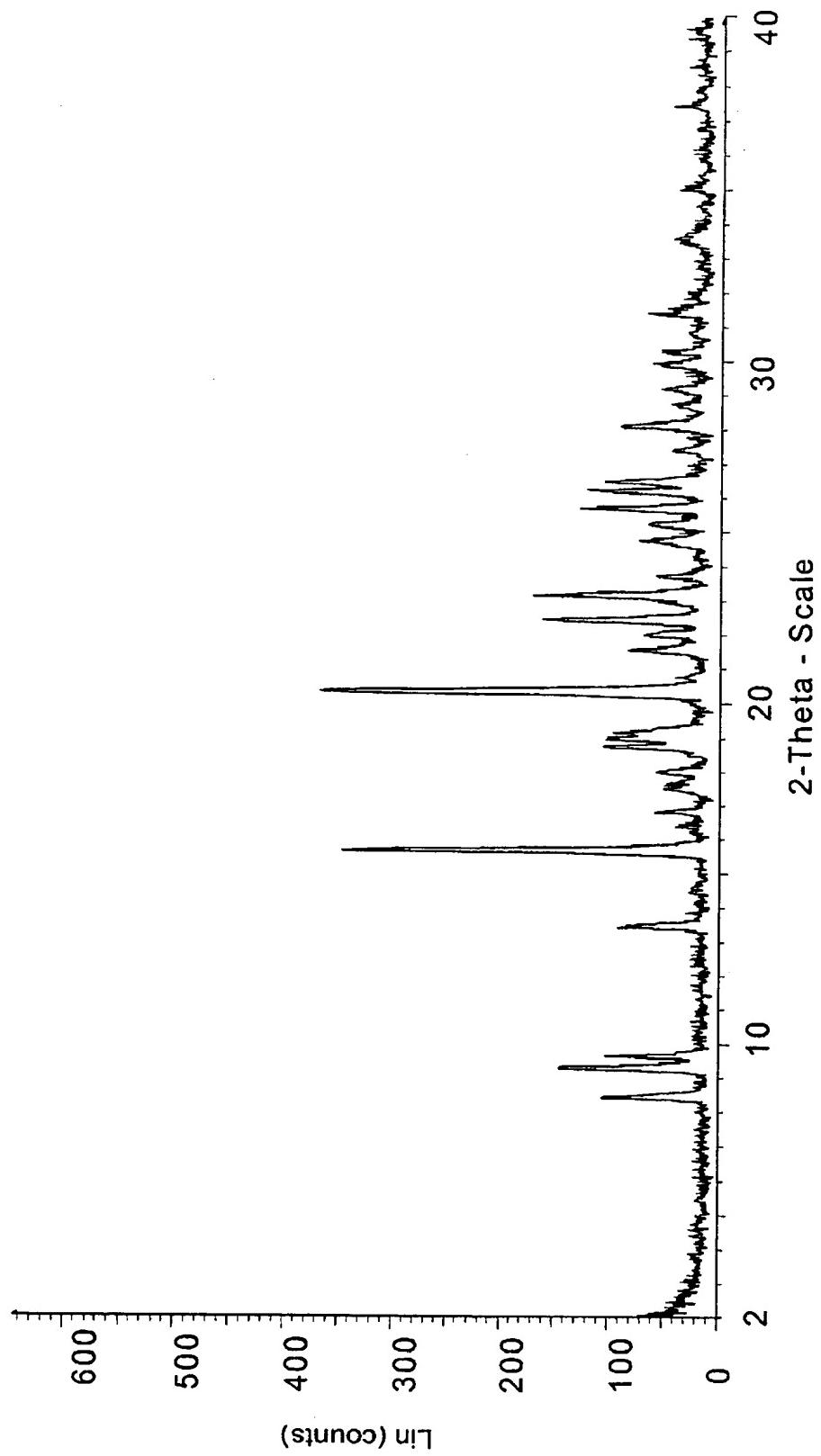
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14. A compound according to Claim 1 which is 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide in crystalline form.

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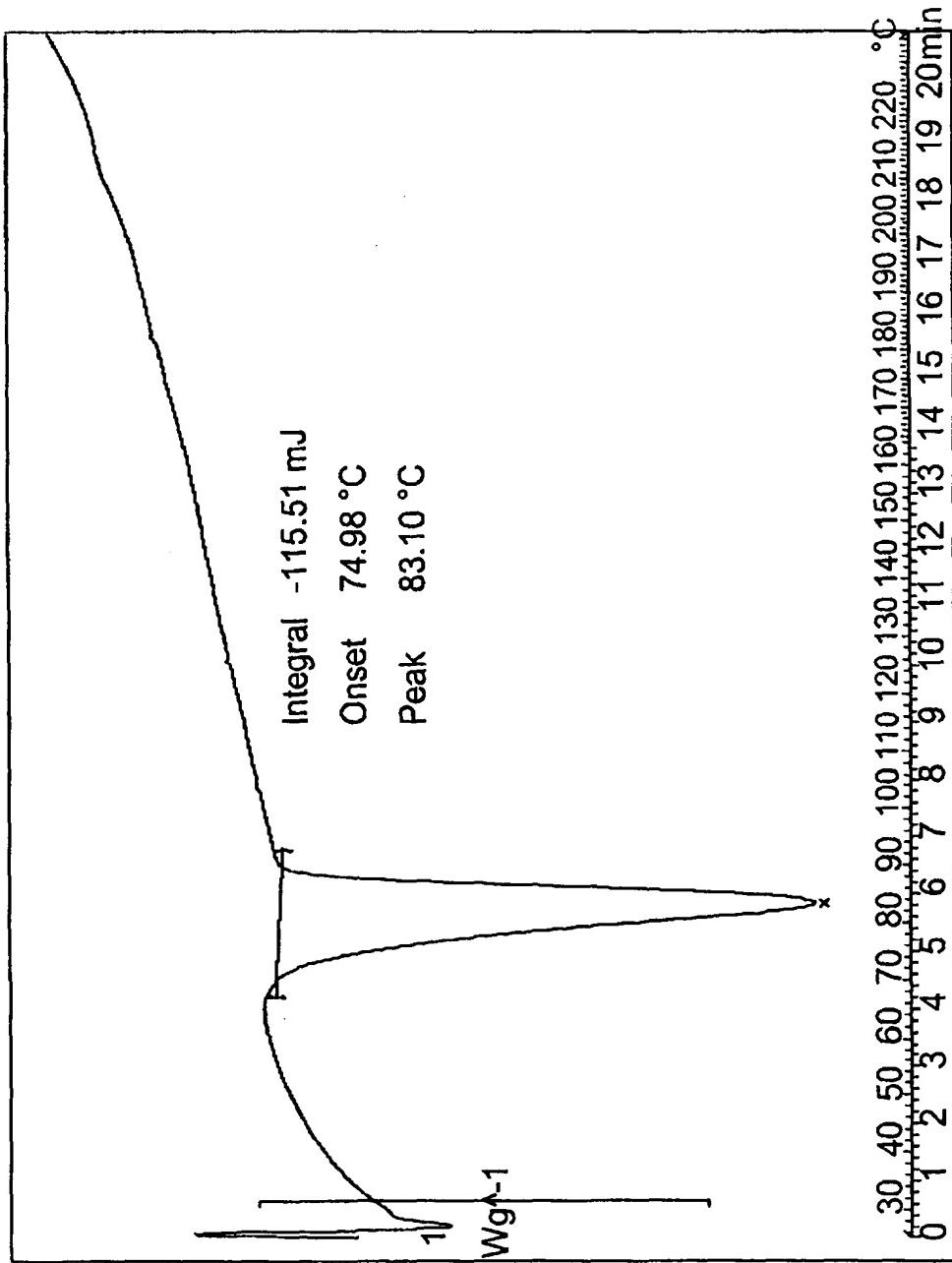
15. A compound according to Claim 14 having an X-ray powder diffraction pattern with at least two specific peaks at about $2\text{-theta} = 20.3^\circ$ and 15.6° when measured using CuKa radiation.

Figure A: X-Ray Powder Diffraction Pattern 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide Form A



2 of 2

Figure B: DSC Thermogram 5-[3-[(2S)-1-(difluoromethoxy)propan-2-yl]oxy-5-[(5-methylpyrazin-2-yl)carbamoyl]phenoxy]-N,N-dimethyl-pyrazine-2-carboxamide Form A



INTERNATIONAL SEARCH REPORT

International application No

PCT/GB2007/004057

A. CLASSIFICATION OF SUBJECT MATTER	INV. C07D241/24 C07D401/12 C07D403/12 A61K31/4439 A61K31/497 A61P3/00
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According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
C07D A61K A61P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, CHEM ABS Data, BEILSTEIN Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 03/015774 A (ASTRAZENECA AB [SE]; ASTRAZENECA UK LTD [GB]; BOYD SCOTT [GB]; CAULKET) 27 February 2003 (2003-02-27) cited in the application page 1, lines 1-10 claims 1-16 -----	1-15
X	EP 1 600 442 A (BANYU PHARMA CO LTD [JP]) 30 November 2005 (2005-11-30) cited in the application claims 1-32 ----- -/-	1-15

Further documents are listed in the continuation of Box C.

See patent family annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *&* document member of the same patent family

Date of the actual completion of the international search

21 January 2008

Date of mailing of the international search report

04/02/2008

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Marzi, Elena

INTERNATIONAL SEARCH REPORT

International application No

PCT/GB2007/004057

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, X	WO 2007/007041 A (ASTRAZENECA AB [SE]; ASTRAZENECA UK LTD [GB]; MCKERRECHER DARREN [GB];) 18 January 2007 (2007-01-18) page 4, lines 4-16 page 117; example 16 page 118; example 16a page 123; example 18 claims 1-16 -----	1-15

INTERNATIONAL SEARCH REPORT

International application No.
PCT/GB2007/004057

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: —
because they relate to subject matter not required to be searched by this Authority, namely:
see FURTHER INFORMATION sheet PCT/ISA/210
2. Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful International search can be carried out, specifically:
3. Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fees, this Authority did not invite payment of additional fees.
3. As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.
- The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
- No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box II.1

Although claims 9-10 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.

Continuation of Box II.1

Claims Nos.: -

Rule 39.1(iv) PCT - Method for treatment of the human or animal body by therapy

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/GB2007/004057

Patent document cited in search report		Publication date		Patent family member(s)		Publication date
WO 03015774	A	27-02-2003		AT 323487 T AT 359781 T AT 334678 T BR 0212008 A CA 2457410 A1 CN 1568185 A DE 60210782 T2 DE 60213678 T2 DE 60219698 T2 DK 1568367 T3 EP 1420784 A1 ES 2285341 T3 ES 2270263 T3 HK 1064598 A1 HU 0401213 A2 IS 7150 A JP 3987829 B2 JP 2005525291 T JP 2005320343 A MX PA04001500 A NO 20040686 A NZ 531193 A US 2005080106 A1 ZA 200401015 A		15-05-2006 15-05-2007 15-08-2006 28-09-2004 27-02-2003 19-01-2005 10-05-2007 09-08-2007 27-12-2007 30-07-2007 26-05-2004 16-11-2007 01-04-2007 29-09-2006 28-12-2004 13-02-2004 10-10-2007 25-08-2005 17-11-2005 14-05-2004 17-02-2004 29-07-2005 14-04-2005 06-05-2005
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WO 2007007041	A	18-01-2007		AR 055074 A1 UY 29656 A1		01-08-2007 28-02-2007